

GenCore version 4.5  
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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:02:40 ; Search time 15.58 Seconds

(Without alignments)  
740.893 Million cell updates/sec

Title: US-09-441-654a-1

Perfect score: 948

Sequence: 1 ADPRRSIHDFCLVSKVYVGR.....ACMLRCFRQENPPLIGSK 170

Scoring table:

BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 195891 seqs, 67900655 residues

Total number of hits satisfying chosen parameters: 195891

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database :

1: pir1:\*  
2: pir2:\*  
3: pir3:\*  
4: pir4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	658	69.4	252	2 JG0185	hepatocyte growth
2	250.5	26.4	1558	2 T34384	hypothetical prote
3	250.5	26.4	2167	2 T34395	hypothetical prote
4	249.5	26.3	352	1 T1B0B1	alpha-1-microglobu
5	246.5	26.0	302	1 T1RTGK	tissue factor path
6	244.5	25.8	123	2 A29652	inter-alpha-trypsi
7	244.5	25.8	352	1 HCHU	alpha-1-microglobu
8	244	25.7	299	2 T46937	tissue factor path
9	243	25.6	300	2 SI2143	lipoprotein-associ
10	242.5	25.6	337	1 T1PGBI	alpha-1-microglobu
11	241.5	25.5	125	1 T1H0B1	alpha-1-microglobu
12	240	25.3	2225	2 T26063	hypothetical prote
13	239.5	25.3	396	2 S53325	tissue factor path
14	237.5	25.1	349	2 T21089	alpha-1-microglobu
15	236.5	24.9	304	1 T22264	tissue factor path
16	235.5	24.8	389	2 S35708	alpha-1-microglobu
17	233.5	24.6	304	1 T1H0GK	tissue factor path
18	222	23.4	1043	1 T19734	hypothetical prote
19	219	23.1	922	2 T23573	hypothetical prote
20	214.5	22.6	235	2 A54951	tissue factor path
21	210	22.2	765	2 T24880	amyloid precursor
22	209	22.0	1743	2 T26859	hypothetical prote
23	208	21.9	751	2 A49974	beta-amyloid precu
24	200	21.1	763	2 A49321	amyloid beta (A4)
25	200	21.1	1599	2 T16210	hypothetical prote
26	195	20.6	1391	2 T20406	hypothetical prote
27	193.5	20.4	111	2 T41082	amyloid precursor
28	193	20.4	2844	2 S28291	hypothetical prote
29	186.5	19.7	747	2 JH0773	Alzheimer's disease

30	186	19.6	484	4 A32761	hypothetical prote
31	186	19.6	770	1 QRH0A4	Alzheimer's disease
32	185.5	19.6	1203	2 T21275	hypothetical prote
33	181.5	19.1	355	1 S22181	gamma-1-microglobu
34	178	18.8	1965	2 T33216	hypothetical prote
35	175.5	18.5	76	2 S03607	Alzheimer's disease
36	174.5	18.4	76	2 S04855	Alzheimer's disease
37	174.5	18.4	76	2 S06678	Alzheimer's disease
38	174.5	18.4	100	2 A32282	Alzheimer's disease
39	174.5	18.4	692	2 T32980	Alzheimer's disease
40	167	17.6	62	2 S07451	hypothetical prote
41	164	17.3	838	2 T20125	hypothetical prote
42	163.5	17.2	372	2 J02556	alpha-1-microglobu
43	161	17.0	1208	2 T27822	hypothetical prote
44	159	16.8	265	2 A53350	Kunitz-type protei
45	156	16.5	64	2 S41399	Kunitz-type protei

## ALIGNMENTS

RESULT 1

JG0185 hepatocyte growth factor activator inhibitor type 2 - mouse

C:Species: Mus musculus (house mouse)

C>Date: 23-Jul-1999 #sequence\_revision 23-Jul-1999 #text\_change 11-May-2000

C:Accession: JG0185

R:Itch, H.; Kataoka, H.; Hamasuna, R.; Kitamura, N.; Koono, M.

Biochem. Biophys. Res. Commun. 255, 740-748, 1999

A:Title: Hepatocyte growth factor activator inhibitor type 2 lacking the first kunitz

A:Reference number: JG0185; PMID:99160423

A:Accession: JG0185

A:Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-252 <110>

A:Cross-references: GB:AF099016

A:Superfamily: animal Kunitz-type proteinase inhibitor homology

F:133-183/Domain: animal Kunitz-type proteinase inhibitor homology <BPI>

Query Match

Best Local Similarity 69.4%; Score 658; DB 2; Length 252;

Matches 116; Conservative 20; Mismatches 34; Indels 0; Gaps 0;

QY	1	ADPRRSIHDFCLVSKVYVGRASAPRMWYVDTGSCQLFYGGCDGNSNVLKEELKK	60
DB	28	ASRELDVHSCGVSKYVGRASIPRMWYVDTGSCQPFYGGCEGNGNYSKEELCK	87
QY	61	CATVENATGDLATSRNADSVSPAPRRDSDHSDMFNFEYCTANAVTGPCRASFP	120
DB	88	CAGVENVETTDNARNRNGADSSVLSVPRKQSAEDLSAEIFNFEYCVKRAVTPCRAPF	147
QY	121	RMYEDERNSCNNEITGCGRKNKNSRSEACMLRFRQENPPLIGSK	170
DB	148	RMYETDKNSCSFTYGGCRKNKNSYLSEACMQHSGRQMHFPLTPGLK	197

RESULT 2

T34394 hypothetical protein C37C3.6a - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C>Date: 29-Oct-1999 #sequence\_revision 29-Oct-1999 #text\_change 29-Oct-1999

C:Accession: T34394

R:Geisel, C.; Bradshaw, H.

submitted to the EMBL Data Library, July 1996

A:Description: The sequence of C. elegans cosmid C37C3.

A:Reference number: Z21518

A:Accession: T34394

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-1558 <GB1>

A:Cross-references: EMBL:U64857; PIDN:AAC25867.1; GSPDB:GN00023; CESP:C37C3.6-

A:Experimental source: strain Bristol N2; clone C37C3

A;Title: Bovine alpha(1)-microglobulin/oligomer. Isolation  
A;Reference number: S68149; MUID:96201710

A;Residues: 1-302 <ENU>

A:Cross-references: DDBJ:D10926; NID:g220916; PIDN:BA01724.1; PID:g220917  
 A:Experimental source: liver  
 C:Comment: This serine proteinase inhibitor regulates clotting by factor Xa-dependent in

C:Superfamily: The first Kunitz-type domain binds the factor VIIa/tissue factor complex; the  
 C:Keywords: anticoagulant; blood coagulation; duplication; glycoprotein; heparin binding  
 F:1-28/Domain: signal sequence #status predicted <SIG>  
 F:29-302/Product: tissue factor pathway inhibitor #status predicted <SIG>  
 F:53-103/Domain: animal Kunitz-type proteinase inhibitor #status predicted <MAT>  
 F:124-174/Domain: animal Kunitz-type proteinase inhibitor homology <BP1>  
 F:222-272/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>  
 F:268-291/Region: heparin binding #status predicted  
 F:53-103/62-86-78-99,124-174,133-157,149-170,222-272,231-255,247-268/Disulfide bonds: #  
 F:63/Inhibitory site: Lys (coagulation factor VII/tissue factor complex) #status predict  
 F:134/Inhibitory site: Arg (coagulation factor X) #status predicted  
 F:144,251,261/Binding site: carbonylate (Asn) (covalent) #status predicted  
 F:222/Inhibitory site: Lys (unidentified proteinase) #status predicted

## Query Match

Best Local Similarity 26.0%; Score 246.5; DB 1; Length 302;  
 Matches 54; Conservative 21; Mismatches 65; Indels 19; Gaps 3;

OY 9 DPLVSVVRCRASPMPWYNTDSCQLFYVGGCDGNSNNYTKRECKKCA-TTEN 67  
 DB 122 DPCFLDEDPICRGFTFRFYNNOSKOCFQKYGCGGCGNSNNFTLECCNTECDPVNEY 181  
 OY 68 ATGDLATSR-----NAADSVSPARRDSEHSDMFNYEECTANAVTGPCRA 117  
 DB 182 QKGDYVNTQITVDTTNNVTPQATKASQWDYDGP-----WCLEPDSGLCKA 233  
 OY 118 SPPRWTFEDVRNSCNFFYGGCGKNGKNSYSEACMLRC 156  
 DB 234 SEKRFYNPALGKCRGFNTGCGGNNNNFTTKDCCNRAC 272

## RESULT 6

A29652

Inter-alpha-trypsin inhibitor (BPI type) - sheep (fragment)  
 C:Species: Ovis orientalis aries, Ovis ammon aries (domestic sheep)  
 C:Date: 31-Dec-1988 #sequence\_revision 31-Dec-1988 #text\_change 16-Jul-1999  
 C:Accession: A29652  
 R:Resp, G.: Hochstrasser, K.; Wachter, E.; Reisinger, P.W.M.  
 Biol. Chem. Hoppe-Seyler 368, 727-731, 1987  
 A:Title: The amino-acid sequence of the trypsin-released inhibitor from sheep inter-alpha-  
 sin inhibitor (XI).  
 A:Reference number: A29652; MUID:87299012  
 A:Accession: A29652  
 A:Molecule type: protein  
 A:Residues: 1-123 <RAS>  
 C:Superfamily: protein HC; animal Kunitz-type proteinase inhibitor homology; lipocalin h  
 C:Keywords: serine proteinase inhibitor  
 F:5-55/Domain: animal Kunitz-type proteinase inhibitor homology <BP1>  
 F:61-111/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>

Query Match 25.8%; Score 244.5; DB 2; Length 123;  
 Best Local Similarity 31.8%; Pred. No. 4e-15;  
 Matches 47; Conservative 15; Mismatches 47; Indels 39; Gaps 1;

OY 9 DPLVSVVRCRASPMPWYNTDSCQLFYVGGCDGNSNNYTKRECKKCA-TYENA 68  
 DB 3 DSCQLYSGPCPLGMFRFYNGTSMACETFFYGGCGKNGNNPSEKCLQTRTV--- 58  
 OY 69 TGDLATSRNAADSVSPARRDSEHSDMFNYEECTANAVTGPCRAFPWYDVER 128  
 DB 59 -----OACMLPIVKGPCRAIGELMADAVK 83  
 OY 129 NSCNNTFYGGCGKNGKNSYSEACMLRC 156  
 DB 84 GRCVRFYGGCGKNGNGQFYSGKECKEYC 111

## RESULT 7

HCHU

alpha-1-microglobulin/inter-alpha-trypsin inhibitor precursor - human  
 N:Alternate names: bikunin; complex-forming glycoprotein heterogeneous in charge (HC  
 rich protein  
 C:Contents: alpha-1-microglobulin (protein HC); inter-alpha-trypsin inhibitor  
 C:Species: Homo sapiens (man)  
 C:Date: 15-Oct-1982 #sequence\_revision 30-Jun-1987 #text\_change 04-Feb-2000  
 C:Accession: S13433; S10778; A93642; A90074; A90686; PNO450; B39079; A61520;  
 3217

R:Veit, H.; Gebhard, W.  
 Biol. Chem. Hoppe-Seyler 371, 1185-1196, 1990

A:Title: Structure of the human alpha(1)-microglobulin-bikunin gene.  
 A:Reference number: S13433; MUID:91214554  
 A:Accession: S13433

A>Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-352 <VEIT>

A:Cross-references: EMBL:X54816; NID:g24475; PIDN:CAA38585.1; PID:g825614; EMBL:X54816  
 R:Diarra-Mehpou, M.; Bouguignon, J.; Sesboue, R.; Salier, J.P.; Leveillard, T.; M  
 Eur. J. Biochem. 191, 131-139, 1990

A:Title: Structural analysis of the human inter-alpha-trypsin inhibitor light-chain g  
 A:Reference number: S10778; MUID:90336621  
 A:Accession: S10778

A>Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-202 <DIA>

R:Kammerer, J.F.; Polazzi, J.O.; Kotlick, M.P.  
 Nucleic Acids Res. 14, 7839-7850, 1986

A:Title: The mRNA for a proteinase inhibitor related to the H1-30 domain of inter-  
 A:Reference number: A93642; MUID:87040757  
 A:Accession: A93642

A:Molecule type: mRNA  
 A:Residues: 1-352 <KAU>

A:Cross-references: S13433; NID:g24478; PIDN:CAA28182.1; PID:g24479  
 R:Lopez-Otin, C.; Grubb, A.O.; Mendez, E.  
 Arch. Biochem. Biophys. 228, 544-554, 1984

A:Title: The complete amino acid sequence of human complex-forming glycoprotein  
 A:Reference number: A90074; MUID:84126849  
 A:Accession: A90074

A:Molecule type: protein  
 A:Residues: 20-56,58-202 <LOP>

A:Experimental source: individual with tubular proteinuria  
 A>Note: no evidence of sequence heterogeneity could be found, in spite of presence of

R:Takegi, T.; Takagi, K.; Kawai, T.  
 Biochem. Biophys. Res. Commun. 98, 997-1001, 1981

A:Title: Complete amino acid sequence of human alpha-1-microglobulin.  
 A:Reference number: A90225; MUID:81184038  
 A:Accession: A90225

A:Molecule type: protein  
 A:Residues: 20-47,58-136,138-141,143-144,146-198 <TAK>

A:Experimental source: pooled urine of patients with tubular proteinuria  
 R:Reisinger, P.; Hochstrasser, K.; Albrecht, G.J.; Lempar, K.; Salier, C.  
 Biol. Chem. Hoppe-Seyler 366, 479-483, 1985

A:Title: Human inter-alpha-trypsin inhibitor: localization of the kunitz-type  
 A:Reference number: A90686; MUID:85225968  
 A:Accession: A90686

A:Molecule type: protein  
 A:Residues: 206-290,291-342,344-350 <REI>

R:Altman, F.; Lacour, B.; Strecker, G.; Parvy, P.; Druet, T.; Daudon, M.  
 Biochem. Biophys. Res. Commun. 191, 1158-1165, 1993

A:Title: Molecular characteristics of uronic-acid-rich protein, a strong inhibitor  
 A:Reference number: PNO450; MUID:93221481  
 A:Accession: PNO450

A:Molecule type: protein  
 A:Residues: 206-214,216-271 <ATM>

R:Englind, J.J.; Salvesen, G.; Heft, S.A.; Thøgersen, I.B.; Rutherford, S.; Pizzo, S  
 J. Biol. Chem. 266, 747-751, 1991

A:Title: Chondroitin 4-sulfate covalently cross-links the chains of the human blood p  
 A:Reference number: A39079; MUID:91093267  
 A:Accession: B39079

A:Molecule type: protein  
 A:Residues: 206-225 <ENG1>

R.Chirat, F.; Baldyck, M.; Mizon, C.; Laroui, S.; Sautiere, P.; Mizon, J.  
Int. J. Biochem. 23, 1201-1203, 1991  
A:Title: A chondroitin-sulfate chain is located on serine-10 of the urinary trypsin inhib  
A:Reference number: A61580; MUID:92175157  
A:Accession: A61580  
A:Molecule type: protein  
A:Residues: 214, 'X', 216-222, 'X' <CH1>  
R.McKeenan, W.L.; Sakagami, Y.; Hoshi, H.; McKeenan, K.A.  
J. Biol. Chem. 261, 5378-5383, 1986  
A:Title: Two apparent human endothelial cell growth factors from human hepatoma cells ar  
A:Reference number: A92583; MUID:8618278  
A:Accession: B25604  
A:Molecule type: protein  
A:Residues: 206-214, 'X', 216-230, 'X', 232-239, 'X', 241-248, 'XX', 251-252, 'X', 254 <MK>  
R.Engelild, J.J.; Thøgersen, I.B.; Pizzo, S.V.; Salvesen, G.  
J. Biol. Chem. 264, 15975-15981, 1989  
A:Title: Analysis of inter-alpha-trypsin inhibitor and a novel trypsin inhibitor, pre-al  
A:Reference number: A92736; MUID:89380192  
A:Accession: C34245  
A:Molecule type: protein  
Residues: 206-225 <ENG2>  
Tataboni, C.; Cortese, R.  
Nucleic Acids Res. 14, 6340, 1986  
A:Title: Sequence of a full length cDNA coding for human protein HC (alpha-1-microglobul  
A:Reference number: A25303; MUID:86312901  
A:Accession: A25303  
A:Molecule type: mRNA  
A:Residues: 1-218, 'HW' <TRA>  
A:Note: this mRNA sequence appears to contain errors after residue 218  
R.Calero, M.; Escarban, J.; Grubb, A.; Mendez, E.  
J. Biol. Chem. 269, 384-389, 1994  
A:Title: Location of a novel type of interpolypeptide chain linkage in the human protein  
A:Reference number: A53110; MUID:94103241  
A:Accession: A53110  
A:Molecule type: protein  
A:Residues: 45-57 <CAL1>  
R.Vetr, H.; Koeigler, M.; Gebhard, W.  
FEBS Lett. 245, 137-140, 1989  
A:Title: The domain structure of the inhibitor subunit of human inter-alpha-trypsin inh  
A:Reference number: S03552; MUID:89171290  
A:Accession: S03552  
A:Status: nucleic acid sequence not shown  
A:Status: preliminary  
A:Molecule type: protein  
A:Residues: 206-352 <VET2>  
R.Malki, N.; Baldyck, M.; Maes, P.; Capon, C.; Mizon, C.; Han, K.K.; Tartar, A.; Fourne  
Biol. Chem. Hoppe-Seyler 373, 1009-1018, 1992  
A:Title: The heavy chains of human plasma inter-alpha-trypsin inhibitor: their isolation  
A:Reference number: S28928; MUID:93039735  
A:Accession: S28928  
A:Status: preliminary  
A:Molecule type: protein  
A:Residues: 206-215 <MA1>  
R.Morille, W.; Capon, C.; Baldyck, M.; Sautiere, P.; Kouch, M.; Michalski, C.; Fournet  
Eur. J. Biochem. 221, 881-888, 1994  
A:Title: Chondroitin sulphate covalently cross-links the three polypeptide chains of int  
A:Reference number: S43466; MUID:94229087  
A:Accession: S43466  
A:Status: preliminary  
A:Molecule type: protein  
A:Residues: 206-221 <MOR>  
R.Wisniewski, H.G.; Burgess, W.H.; Oppenheim, J.D.; Vallock, J.  
Biochemistry 33, 7423-7429, 1994  
A:Title: TSG-6, an arthritis-associated hyaluronan binding protein, forms a stable comp  
A:Reference number: A53642; MUID:94271799  
A:Accession: A53642  
A:Status: preliminary  
A:Molecule type: protein  
A:Residues: 206-217 <WIS>  
R.Calero, M.; Mendez, E.; Garcia, E.  
Biochim. Biophys. Acta 1249, 91-99, 1995  
A:Title: Expression of the human complex-forming glycoprotein HC (alpha-1-microglobulin)  
A:Reference number: S55688; MUID:95284116  
A:Accession: S55688

A:Molecule type: protein  
A:Residues: 20-24 <CHL2>  
R.Bourguignon, J.; D'Arra-Mehrpour, M.; Seeboue, R.; Frain, M.; Sala-Trepat, J.M.; Ma  
Biochem. Biophys. Res. Commun. 131, 1146-1153, 1995  
A:Title: Human inter-alpha-trypsin-inhibitor: characterization and partial nucleotide  
A:Reference number: 152208; MUID:86025577  
A:Accession: 152208  
A:Status: translated from GB/EMBL/DBD1  
A:Molecule type: mRNA  
A:Residues: 302-352 <BOU>  
A:Cross-references: GB:M11562; NID:g186587; PIDN:AA59194.1; PID:9307077  
R.Mojcik, E.G.C.; van den Berg, M.; van der Linden, I.K.; Poort, S.R.; Cupers, R.; Pe  
Biochem. J. 311, 753-759, 1995  
A:Title: Factor IX Zúrtphen: a Cys(18) -> Arg mutation results in formation of a heter  
A:Reference number: S59509; MUID:96067589  
A:Accession: S59509  
A:Molecule type: protein  
A:Residues: 27-35, 'Y', 37 <MOU>  
R.Amani, F.; Mizon, J.; Khan, S.R.  
Eur. J. Biochem. 236, 984-990, 1996  
A:Title: Identification of uronic-acid-rich protein as urinary bikunin, the light cha  
A:Reference number: S66434; MUID:96270753  
A:Accession: S66434  
A:Molecule type: protein  
A:Residues: 206-214, 'X', 216-230 <ATM2>  
R.Aksteröom, B.; Bratt, T.; Englund, J.J.  
FEBS Lett. 362, 50-54, 1995  
A:Title: Formation of the alpha(1)-microglobulin chromophore in mammalian and insect  
A:Reference number: S68728; MUID:95212582  
A:Accession: S68728  
A:Molecule type: protein  
A:Residues: 89-100 <AKE>  
R.Jessen, T.E.; Faarvang, K.L.; Ploung, M.  
FEBS Lett. 230, 195-200, 1988  
A:Title: Carbohydrate as covalent crosslink in human inter-alpha-trypsin inhibitor  
A:Reference number: S02431; MUID:88167187  
A:Accession: S02431  
A:Molecule type: protein  
A:Residues: 206-214, 'X', 216-217 <JES>  
R.Lopez, C.; Grubb, A.; Mendez, E.  
FEBS Lett. 144, 349-353, 1982  
A:Title: Human protein HC displays variability in its carboxyl-terminal amino acid se  
A:Reference number: A91304  
A:Contents: annotation: variant of alpha-1-microglobulin  
A:Note: pooled urine samples contained two forms of this protein, both lacking 57-Tyr  
R.Hochstrasser, K.; Schonberger, O.L.; Rosenanth, I.; Wachter, E.  
Hoppe-Seyler's Z. Physiol. Chem. 362, 1357-1362, 1981  
A:Title: Kunitz-type proteinase inhibitors derived by limited proteolysis of the  
by affinity chromatography.  
A:Reference number: A91698; MUID:82074265  
A:Contents: annotation: carbohydrate binding sites  
R.Moril, M.; Travis, J.  
Biol. Chem. Hoppe-Seyler 366, 19-21, 1985  
A:Title: The reactive site of human inter-alpha-trypsin inhibitor is in the 9  
A:Reference number: A90687; MUID:85225940  
A:Contents: annotation: inhibitory site  
A:Note: in vitro, the first twelve residues of the amino end of the inhibitor  
A:Comment: Alpha-1-microglobulin and inter-alpha-trypsin inhibitor are proteolytic  
C:Comment: Alpha-1-microglobulin occurs in many physiological fluids including  
It contains at least one brown-yellow chromophore.

Query Match 25.8%; Score 244.5; DB 1; Length 352;  
Best Local Similarity 32.4%; Pred No.1-2e-14;  
Matches 48; Conservative 14; Mismatches 39; Gaps 1;  
DB 229 DSCQLGYSAGPCMGMTSRFYNGNSMACEFFGCGCMGNMFVEKECLQCTRTVA-- 286  
9 DECLYSKVGCRASMPRMWYNTDSCQLFYTGCGDGSNNYLRKECIKCAATYENA 68  
DB 229 DSCQLGYSAGPCMGMTSRFYNGNSMACEFFGCGCMGNMFVEKECLQCTRTVA-- 286  
QY 69 TGDLTATSNALDSSVPAPRROSESDHSSDMFEYEEYCANNAVTCGRASPRFWYDVER 128  
DB 287 -----CNLPYVRCRAFIQLMAFDVAK 309



QY 129 NSCNFIYGGCGKGNKSYSEACMLRC 156  
 Db 310 GKCVLFPGGCGGNKFKYSKECEKRC 337

# RESULT 8

Tissue factor pathway inhibitor - rabbit  
 C:Species: Oryctolagus cuniculus (domestic rabbit)  
 C:Date: 04-Sep-1997 #sequence\_revision 04-Sep-1997 #text\_change 13-Aug-1999  
 C:Accession: I46937  
 R:Belasquez, A.; Kumpuswamy, M.N.; Birktoft, J.J.; Bajaj, S.P.  
 Thromb. Res. 69, 547-553, 1993  
 A:Title: Revised cDNA sequence of rabbit tissue factor pathway inhibitor.  
 A:Reference number: I46937; MUID:93276427  
 A:Accession: I46937  
 A:Status: preliminary  
 A:Molecule type: mRNA  
 A:Residues: 1-299 <BEL>

Cross-references: GB:S61902; NID:q386015; PIDN:AA626836.1; PID:q386016  
 Superfamily: tissue factor pathway inhibitor; animal Kunitz-type proteinase inhibitor  
 F:49-99/Domain: animal Kunitz-type proteinase inhibitor homology <BP1>  
 F:120-170/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>  
 F:212-262/Domain: animal Kunitz-type proteinase inhibitor homology <BP12>

Query Match Best Local Similarity 25.7%; Score 244; DB 2; Length 299;  
 Matches 48; Conservative 23; Mismatches 61; Indels 28; Gaps 2;

QY 4 ERSIHDFCLVSKVGRASMPRMWYNTDSCOLFVYGGCGGNSNNYLTKECLKCAT 63  
 Db 42 QKPTHSFCAMKVDGDCRCAYIKRFNNILAHQCEEFYGGCGENRFSLECKEKCAR 101  
 QY 64 VTEAATGDLATSRNAADSVSPAPRRDSEDDHSSDMENYECYANAVTGPCRASPPRM 123  
 Db 102 DYPMATTKLTFQKGRPD-----FCFLDEDPGICRQYITRYF 137  
 QY 124 FDEERNSCNNFIYGGCGKGNKSYSEACMLRCFROENP 163  
 Db 138 YNNSKQCEKRFKYGCGCLGNLFESLECKKNTC---ENP 173

# RESULT 9

liporotein-associated coagulation inhibitor precursor - rabbit  
 A:Alternate names: endothelial cell coagulation inhibitor; endothelial cell tissue facto

C:Species: Oryctolagus cuniculus (domestic rabbit)  
 C:Date: 19-Mar-1997 #sequence\_revision 19-Mar-1997 #text\_change 16-Jul-1999  
 C:Accession: S12143; A61373

R:Wesselschmidt, R.L.; Girard, T.J.; Broze Jr., G.J.  
 Nucleic Acids Res. 18, 6440, 1990  
 A:Title: cDNA sequence of rabbit liporotein-associated coagulation inhibitor.  
 A:Reference number: S12143; MUID:91057146  
 A:Accession: S12143

A:Status: preliminary  
 A:Molecule type: protein  
 A:Residues: 1-300 <WES>

A:Cross-references: EMBL:X54708; NID:q1612; PIDN:CAA8515.1; PID:q1613  
 R:Colburn, P.; Grab, J.W.; Buonassisi, V.  
 J. Cell. Physiol. 148, 320-326, 1991  
 A:Title: Enhanced inhibition of tissue factor by the extended form of an endothelial cell

A:Reference number: A61373; MUID:91349227  
 A:Accession: A61373

A:Molecule type: protein  
 A:Residues: 25-33, 'X', '35-46 <COL>  
 C:Superfamily: tissue factor pathway inhibitor; animal Kunitz-type proteinase inhibitor  
 C:Keywords: anticoagulant; glycoprotein  
 F:50-100/Domain: animal Kunitz-type proteinase inhibitor homology <BP1>  
 F:121-171/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>  
 F:213-263/Domain: animal Kunitz-type proteinase inhibitor homology <BP3>

Query Match Best Local Similarity 25.6%; Score 243; DB 2; Length 300;  
 Matches 48; Conservative 23; Mismatches 61; Indels 28; Gaps 2;

QY 4 ERSIHDFCLVSKVGRASMPRMWYNTDSCOLFVYGGCGGNSNNYLTKECLKCAT 63  
 Db 43 QKPTHSFCAMKVDGDCRCAYIKRFNNILAHQCEEFYGGCGENRFSLECKEKCAR 102  
 QY 64 VTEAATGDLATSRNAADSVSPAPRRDSEDDHSSDMENYECYANAVTGPCRASPPRM 123  
 Db 103 DYPMATTKLTFQKGRPD-----FCFLDEDPGICRQYITRYF 138  
 QY 124 FDEERNSCNNFIYGGCGKGNKSYSEACMLRCFROENP 163  
 Db 139 YNNSKQCEKRFKYGCGCLGNLFESLECKKNTC---ENP 174

# RESULT 10

alpha-1-microglobulin/inter-alpha-trypsin inhibitor precursor - pig (fragment)  
 A:Alternate names: bikunin; ITI; PI-14 (inhibitory fragment of ITI)  
 C:Species: Sus scrofa domestica (domestic pig)  
 C:Date: 30-Jun-1987 #sequence\_revision 04-Feb-2000 #text\_change 04-Feb-2000  
 R:Gebhard, W.; Schreimüller, T.; Vetr, H.; Wächter, E.; Hochstrasser, K.  
 FEBS Lett. 269, 32-36, 1990  
 A:Title: Complementary DNA and deduced amino acid sequences of porcine alpha1-microgl  
 A:Reference number: S11066; MUID:90353595  
 A:Accession: S11066

A:Status: preliminary  
 A:Molecule type: mRNA  
 A:Residues: 1-337 <GEB>  
 A:Cross-references: EMBL:X53685; NID:q1877; PIDN:CAA37725.1; PID:q1878  
 R:Tavakoli, A.  
 Biochim. Biophys. Acta 1088, 47-56, 1991

A:Title: Molecular cloning of porcine alpha(1)-microglobulin/Hi-30 reveals developmen  
 A:Reference number: S13493; MUID:91113729  
 A:Accession: S13493

A:Status: preliminary  
 A:Molecule type: mRNA  
 A:Residues: 'M', '3-48', 'M', '50-337 <TAV>  
 A:Cross-references: GB:X52087; NID:q1881; PIDN:CAA6306.1; PID:q1882  
 A:Note: the authors translated the codon GTC for residue 2 as a Met initiation codon  
 R:Hochstrasser, K.; Wächter, E.; Albrecht, G.J.; Reisinger, P.  
 Biol. Chem. Hoppe-Seyler 366, 473-478, 1985

A:Title: Kunitz-type proteinase inhibitors derived by limited proteolysis of the inte  
 A:Reference number: A90685; MUID:85225967  
 A:Accession: A01208

A:Molecule type: protein  
 A:Residues: 212-258, 'Q', '260-269', 'S', '271-277', 'Q', '279-282', 'A', '284', 'IR', '287-292', 'A', '294-  
 C:Comment: This inhibitory fragment, released from native ITI after limited proteolys  
 C:Comment: The amino acid at position P2 (228-Met) appears to determine the specific  
 nd elastase; those with leucine interact strongly.  
 C:Superfamily: protein HC; animal Kunitz-type proteinase inhibitor homology; lipo

C:Keywords: duplication; glycoprotein; plasma; serine proteinase inhibitor  
 F:20-173/Domain: lipoalain homology <LIP>  
 F:216-266/Domain: animal Kunitz-type proteinase inhibitor homology <BP1>  
 F:272-322/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>  
 F:216-266, 225-249, 241-262, 272-322, 281-305, 297-318/Disulfide bonds: #status pre  
 F:226/Inhibitory site: Leu (chymotrypsin, elastase) #status predicted  
 F:235/Binding site: carbonylate (Asn) (covalent) #status experimental  
 F:282/Inhibitory site: Arg (trypsin) #status predicted

Query Match Best Local Similarity 25.6%; Score 242.5; DB 1; Length 337;  
 Matches 47; Conservative 19; Mismatches 43; Indels 39; Gaps

QY 9 DECLVSKVGRASMPRMWYNTDSCOLFVYGGCGGNSNNYLTKECLKCAVTEENA 68  
 Db 214 DSCQLGYSQGPCLGMKRFYNGSSMACETFRYGGCGGNFVSEKCLQCRIV--- 269

QY 69 TGLATSRNADSSVSPAPRRDSEHSDMFNEEYECTANAVTGPCRASFPRMVFVER 128  
 DB 270 -----EACSLPIYSGPCRGFFQJMAFDAAVQ 294  
 QY 129 NSCNFFIYGGCRGNKNSYRSEACMLRC 156  
 DB 295 GKCVLFYGGCGNGNGNFYSERCKEYEC 322

## RESULT 11

TIRHOB1  
 alpha-1-microglobulin/inter-alpha-trypsin inhibitor - horse (fragment)  
 N:Alternate names: EI-14 (inhibitory fragment of IRI); IRI; trypsin inhibitor, E-UTI  
 C:Species: Equus caballus (domestic horse)  
 C>Date: 30-Jun-1987 #sequence\_revision 04-Feb-2000 #text\_change 05-May-2000  
 C:Accession: A01210; A45653  
 R:Hochstrasser, K.; Wachter, E.; Albrecht, G.J.; Reisinger, P.  
 Biol. Chem. Hoppe-Seyler 366, 473-478, 1985  
 Title: Kunitz-type proteinase inhibitors derived by limited proteolysis of the inter-  
 Reference number: A90685; MUID:85225967  
 A:Accession: A01210  
 A:Molecule type: protein  
 A:Residues: 3-125 <HOC>  
 R:Veeraratnam, K.; Singh, K.; Wachter, E.; Hochstrasser, K.  
 Biochem. Int. 26, 405-413, 1992  
 A:Title: Characterization of a trypsin inhibitor from equine urine.  
 A:Reference number: A45653; MUID:92328613  
 A:Accession: A45653  
 A:Status: preliminary  
 A:Molecule type: protein  
 A:Residues: 1-12, E', 14-33 <VEE>  
 A:Cross-references: PIDN:AB22430.1; PID:g250858  
 A:Experimental source: urine  
 A:Note: sequence extracted from NCBI backbone (NCBI:107966)  
 C:Comment: This inhibitory fragment, released from native IRI after limited proteolysis  
 first domain interacts weakly with PMN-granulocytic elastase and not at all with pancrea  
 C:Comment: The amino acid at position P2' (19-Met) appears to determine the specificity  
 d elastase: those with leucine interact strongly.  
 C:Keywords: duplication; glycoprotein; plasma; serine proteinase inhibitor  
 C:Keywords: duplication; glycoprotein; plasma; serine proteinase inhibitor  
 F:7-57/Domain: animal Kunitz-type proteinase inhibitor homology <BP1>  
 F:63-113/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>  
 F:57-16-40,32-53,63-113,72-96,88-109/Distal bonds: #status predicted  
 F:17/inhibitory site: Leu (chymotrypsin, elastase) #status predicted  
 F:26/Binding site: carbonylate (Asn) (covalent) #status experimental  
 F:73/inhibitory site: Arg (trypsin) #status predicted

Query Match 25.5%; Score 241.5; DB 1; Length 125;  
 Best Local Similarity 31.8%; Pred. No. 7.5e-15;  
 Matches 47; Conservative 15; Mismatches 47; Indels 39; Gaps 1;  
 QY 9 DFCVSKVYVGRASMPRMWYNTDSCQLFYVGGCDGNSNNYLTKECLKCAATVENA 68  
 DB 5 DSCQDHAQGCCLGMSIYFNGISMACETFOYGGCLGNGNFSQKCELOTCRTVA-- 62  
 QY 69 TGLATSRNADSSVSPAPRRDSEHSDMFNEEYECTANAVTGPCRASFPRMVFVER 128  
 DB 63 -----CNLPYVQPCAFIRLMAFDAAQ 85  
 QY 129 NSCNFFIYGGCRGNKNSYRSEACMLRC 156  
 DB 86 GKCVLFYGGCGNGNGNFYSERCKEYEC 113

RESULT 12  
 T26063  
 hypothetical protein W01F3.3 - Caenorhabditis elegans  
 C:Species: Caenorhabditis elegans  
 C>Date: 15-Oct-1999 #sequence\_revision 15-Oct-1999 #text\_change 15-Oct-1999  
 C:Accession: T26063  
 R:Cummings, P.

submitted to the EMBL Data Library, March 1997

A:Reference number: 220145  
 A:Accession: T26063  
 A:Status: preliminary; translated from GB/EMBL/DBJ  
 A:Molecule type: DNA  
 A:Residues: 1-2225 <WIL>  
 A:Cross-references: EMBL:292815; PIDN:CAB07294.1; GSPDB:GN00023; CESP:W01F3.3  
 A:Experimental source: clone W01F3  
 C:Genetics:  
 A:Gene: CESP:W01F3.3  
 A:Map position: 5  
 A:Introns: 33/1; 56/1; 100/1; 142/3; 271/3; 451/1; 525/3; 774/1; 1093/1; 1176/1;

Query Match 25.3%; Score 240; DB 2; Length 2225;  
 Best Local Similarity 29.3%; Pred. No. 2.1e-13;  
 Matches 46; Conservative 21; Mismatches 58; Indels 32; Gaps 2;  
 QY 11 CLVSKVYVGRASMPRMWYNTDSCQLFYVGGCDGNSNNYLTKECLKCAATVENA 69  
 DB 777 CLHRDSCNGCGVRFNFDEKKNCDVFTYTCGQGNNGNFASKECMATCKRKEPTPSA 83;  
 QY 69 TGLATSRNADSSVSPAPRRDSEHSDMFNEEYECTANAVTGPCRASFPRMVFVER 128  
 DB 837 TPD-----FSQVCSNDYDAGECNGVFERAFDALA 663  
 QY 129 NSCNFFIYGGCRGNKNSYRSEACMLRCFROENPPL 165  
 DB 867 ODCRAFYGGCGNGNGNFATMQECSRVMAMKSPV 903

## RESULT 13

S53325  
 tissue factor pathway inhibitor - rabbit  
 C:Species: Oryctolagus cuniculus (domestic rabbit)  
 C>Date: 01-Aug-1995 #sequence\_revision 01-Sep-1995 #text\_change 16-Jul-1999  
 C:Accession: S53325  
 R:Girard, J. J.; Gailani, D.; Broze Jr., G.J.  
 Biochem. J. 303, 923-928, 1994  
 A:Title: Complementary DNA sequencing of canine tissue factor pathway inhibitor revea  
 A:Reference number: S53325; MUID:95071310  
 A:Accession: S53325  
 A:Status: preliminary  
 A:Molecule type: mRNA  
 A:Residues: 1-336 <GIR>  
 C:Keywords: serine proteinase inhibitor  
 C:Superfamily: animal Kunitz-type proteinase inhibitor homology  
 F:53-103/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>  
 F:125-175/Domain: animal Kunitz-type proteinase inhibitor homology <BP3>  
 F:309-359/Domain: animal Kunitz-type proteinase inhibitor homology

Query Match 25.3%; Score 239.5; DB 2; Length 396;  
 Best Local Similarity 29.6%; Pred. No. 3.8e-14;  
 Matches 45; Conservative 24; Mismatches 60; Indels 23; Gaps 1;  
 QY 5 RSHDPLVSKVYVGRASMPRMWYNTDSCQLFYVGGCDGNSNNYLTKECLKCAATV 64  
 DB 47 RLHSFALRADNGPCRAMIRNFNFHITQCEEFYVGGCEGNGNRFESIECEKCVRV 106  
 QY 65 TENATGLATSRNADSSVSPAPRRDSEHSDMFNEEYECTANAVTGPCRASFPRMVF 124  
 DB 107 YPKA-----KTELEKLEKPDCHANNEDSGLCRGVTRYY 143  
 QY 125 DYERNSCNFFIYGGCRGNKNSYRSEACMLRC 156  
 DB 144 NVSSKCEGFKYGGCLGGLNLFETLEQCKNTC 175

RESULT 14  
 S51089  
 alpha-1-microglobulin/inter-alpha-trypsin inhibitor light chain precursor - rat  
 N:Alternate names: acid-stable proteinase inhibitor; bikunin; tryptastin

```

RESULT 15
JC2264
tissue factor pathway inhibitor precursor - rhesus macaque
N:Alternate names: extrinsic pathway inhibitor; lipoprotein-associated coagulation
C:Species: Macaca mulatta (rhesus macaque)
C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 21-Jul-2000
C:Accession: J02264
R:Kamei, S.; Kamikubo, Y.; Hamuro, T.; Fujimoto, H.; Ishihara, M.; Yonemura, H.; Miyazaki,
J. Blochem. 115, 708-714, 1994
A:Title: Amino acid sequence and inhibitory activity of rhesus monkey tissue factor
A:Reference number: J02264; MUID:94375417
A:Accession: J02264
A:Molecule type: mRNA
A:Residues: 1-304 <KAM>
A:Cross-references: GB:S7337; NID:9685016; PIDN:AB031955.1; PID:9665017
A:Experimental source: liver
C:Comment: This protein inhibits the activities of factor Xa and tissue factor-
C:superfamily: tissue factor pathway inhibitor; animal Kunitz-type proteinase
C:Keywords: anticoagulant; glycoprotein; serine proteinase inhibitor
F:1-28/Domain: signal sequence #status predicted <SIG>
F:29-304/Product: tissue factor pathway inhibitor #status predicted <NAT>
F:54-104/Domain: animal Kunitz-type proteinase inhibitor homology <BP1>
F:125-175/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>
F:217-267/Domain: animal Kunitz-type proteinase inhibitor homology <BP3>
F:54-104,63-87,79-100,125-175,134-158,150-171,217-267,226-250,242-263/Disulfide bonds
F:634/inhibitory site: lys (coagulation factor VII/tissue factor complex) #status pred
F:135/inhibitory site: Arg (coagulation factor X) #status predicted
F:145,195,256/binding site: Arg (carbohydrate (Asn) (covalent) #status predicted
F:227/inhibitory site: Arg (unidentified proteinase) #status predicted

```

```
Search completed: January 31, 2001, 15:03:23
Job time: 43 sec
```

[illegible]

\_\_\_\_\_

GenCore version 4.5  
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 31, 2001, 15:03:08 ; Search time 10.15 Seconds

(without alignments)  
540.886 Million cell updates/sec

Title: US-09-441-654A-1

Perfect score: 948

Sequence: 1 ADERSIHDFCLVSKVGRG.....ACMLRCFRQENPPLGSK 170

Scoring table:

BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 88757 seqs, 32294092 residues

Minimum number of hits satisfying chosen parameters: 88757

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_39:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	249.5	26.3	352	1	AMBP_BOVIN
2	247.5	26.1	346	1	AMBP_MERON
3	246.5	26.0	302	1	TFPI_RAT
4	244.5	25.8	123	1	IATR_SHEEP
5	244.5	25.8	352	1	AMBP_HUMAN
6	244.5	25.7	300	1	TFPI_RABT
7	242.5	25.6	337	1	AMBP_PIG
8	241.5	25.5	123	1	IATR_HORSE
9	237.5	25.1	349	1	AMBP_RAT
10	236.5	24.9	304	1	TFPI_MACMU
11	235.5	24.8	349	1	AMBP_MESAU
12	235.5	24.8	349	1	AMBP_MOUSE
13	233.5	24.6	304	1	TFPI_HUMAN
14	214.5	22.2	765	1	TFPI_HUMAN
15	210	22.2	765	1	TFPI_RAT
16	200	21.1	763	1	TFPI_RAT
17	191	20.1	1416	1	AMBP_HUMAN
18	187.5	19.8	770	1	YNB1_CAMEL
19	186	19.6	751	1	A4_RAT
20	186	19.6	770	1	A4_SAISC
21	183.5	19.4	770	1	A4_HUMAN
22	181.5	19.1	355	1	AMBP_PIEPL
23	175.5	18.5	69	1	CRPT_BOOMT
24	174.5	18.4	76	1	A4_MACMU
25	174.5	18.4	76	1	A4_MACMU
26	167	17.6	62	1	IP52_ANESU
27	159	16.8	265	1	TKD1_SHEEP
28	156	16.5	64	1	TKD1_HUMAN
29	155	16.5	164	1	TKD1_BOVIN
30	155	16.4	60	1	IBP8_BOVIN
31	153	16.1	100	1	IBP1_TACTR
32	153	16.1	100	1	IBP1_BOVIN
33	152.5	16.1	100	1	IBP1_BOVIN

34	152	16.0	65	1	IVB3_VIPAA	P00992 vipera ammoo
35	150.5	15.9	60	1	IVB2_DABRU	P00990 dabola russ
36	150.5	15.9	122	1	UPT1_PIG	P029100 sus scrofa
37	150	15.8	110	1	IBP_PIG	P00993 carotis rir
38	149	15.7	61	1	IBP_CARCR	P00994 vipera ammoo
39	147	15.5	58	1	ISIR_HELPO	helix cornu
40	147	15.5	67	1	IBPC_BOVIN	P00976 bos taurus
41	145	15.3	55	1	ISH2_STOHE	P8129 stochellic
42	145	15.3	62	1	IVB2_ERMA	P24541 eristocopa
43	143	15.1	83	1	ELAC_MACEU	P02845 macropus cu
44	142	15.0	83	1	CA36_CHICK	P15989 gallus gall
45	141	14.9	65	1	IVB1_BUNFA	P25660 bungarus la

## ALIGNMENTS

RESULT	ID	STANDARD	PRT	352 AA.
1	AMBP_BOVIN			
AC	P00978; P35420; Q28020;			
DT	21-JUL-1986 (Rel. 01, Created)			
DT	01-NOV-1997 (Rel. 35, Last sequence update)			
DT	15-JUL-1999 (Rel. 38, Last annotation update)			
DE	AMBP PROTEIN PRECURSOR (CONTAINS: ALPHA-1-MICROGLOBULIN; INTER-ALPHA-TRYPsin INHIBITOR LIGHT CHAIN (ITI-1C) (BIKUNIN) (HI-30) (BI-14) (CUMULUS EXTRACELLULAR MATRIX STABILIZING FACTOR) (ESF)).			
GN	AMBP OR ITIL.			
OS	Bos taurus (Bovine).			
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;			
CC	Bovidae; Bovinae; Bos.			
RP	[1]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=LIVER;			
RX	MEDLINE=96201710; PubMed=8611630;			
RA	Lindqvist A., Akerstrom B.;			
RT	"Bovine alpha 1-microglobulin/bikunin. Isolation and characterization of liver cDNA and urinary alpha 1-microglobulin.";			
RL	Biochim. Biophys. Acta 1306:98-106(1996).			
RN	[2]			
RN	SEQUENCE OF 227-349.			
RP	MEDLINE=85225967; PubMed=2408637;			
RX	Hochstrasser K., Wachter E., Albrecht G.J., Reisinger P.;			
RA	"Kunitz-type proteinase inhibitors derived by limited proteolysis of the inter-alpha-trypsin inhibitor, X. The amino-acid sequences of the trypsin-released inhibitors from horse and pig inter-alpha-trypsin inhibitors.";			
RL	Biol. Chem. Hoppe-Seyler 366:473-478(1985).			
RN	[3]			
RN	SEQUENCE OF 227-348.			
RP	MEDLINE=84133807; PubMed=6199275;			
RX	Hochstrasser K., Wachter E.;			
RA	"Kunitz-type proteinase inhibitors derived by limited proteolysis of the inter-alpha-trypsin inhibitor, VII. Determination of the amino-acid sequence of the trypsin-released inhibitor from bovine inter-alpha-trypsin inhibitor.";			
RL	Hoppe-Seyler's Z. Physiol. Chem. 364:1679-1687(1983).			
RN	[4]			
RN	SEQUENCE OF 206-219.			
RP	TISSUE=FETAL SERUM;			
RX	MEDLINE=92291130; PubMed=1376324;			
RA	Chen L., Mo S.-J., Larsen W.J.;			
RT	"Identification of a factor in fetal bovine serum that stabilizes the cumulus extracellular matrix. A role for a member of the inter-alpha-trypsin inhibitor family.";			
RL	J. Biol. Chem. 267:12380-12386(1992).			
RN	[5]			
RN	REACTIVE SITES.			
RP	MEDLINE=84133808; PubMed=6199276;			
RX	Hochstrasser K., Albrecht G.J., Schoenberger O.L., Wachter E.;			
RA	"Kunitz-type proteinase inhibitors derived by limited proteolysis of the inter-alpha-trypsin inhibitor, VII. Characterization of the			

RT bovine inhibitor as double-headed trypsin-elastase inhibitor.":  
 RL Hoppe-Seyler's 2. Physiol. Chem. 364:1689-1696(1983).  
 CC -1- FUNCTION: ALPHA-1-MICROGLOBULIN OCCURS IN MANY PHYSIOLOGICAL  
 CC FLUIDS INCLUDING PLASMA, URINE, AND CEREBROSPINAL FLUID. IT  
 CC APPEARS NOT ONLY AS A FREE MONOMER BUT ALSO IN COMPLEXES WITH IGA  
 CC AND ALBUMIN.  
 CC -1- FUNCTION: INTER-ALPHA-TRYPsin INHIBITOR, PRESENT IN PLASMA AND  
 CC URINE, INHIBITS TRYPSIN, PLASMIN, AND LYSOSOMAL GRANULOCYtic  
 CC ELASTASE.  
 CC -1- FUNCTION: MAY DIFFUSE INTO FOLLICULAR FLUID AFTER AN OVULATORY  
 CC STIMULUS TO ACT AS STRUCTURAL LINKER THAT ENSURE NORMAL COMULUS  
 CC EXPANSION, THROUGH STABILIZATION OF THE COMULUS EXTRACELLULAR  
 CC MATRIX THUS SUPPORTING THE PROCESS OF OVULATION.  
 CC -1- PTM: THE PRECURSOR IS PROTEOLYTICALLY PROCESSED INTO TWO  
 CC SEPARATELY FUNCTIONING PROTEINS.  
 CC -1- PTM: HC CONTAINS A COVALENTLY LINKED BROWN-YELLOW CHROMOPHORE.  
 CC -1- SIMILARITY: IN THE N-TERMINAL SECTION, BELONGS TO THE LIPOCALIN  
 CC FAMILY.  
 CC -1- SIMILARITY: IN THE C-TERMINAL SECTION, BELONGS TO THE BPTI/KUNITZ  
 CC FAMILY OF INHIBITORS.  
 CC -----  
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration  
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 CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch))  
 CC -----  
 CC EMBL: U35642; AAB07599.1; -  
 CC PIR: A01209; TIBOBI.  
 CC HSSP: P10646; IADZ.  
 CC INTERPRO: IPR000566; -  
 CC INTERPRO: IPR002223; -  
 CC INTERPRO: IPR002345; -  
 CC PFAM: PF00014; Kunitz\_BPTI; 2.  
 CC PFAM: PF00061; Lipocalin; 1.  
 CC PRINTS: PR00179; LIPOCALIN.  
 CC PRINTS: PR00759; BASICTPASE.  
 CC PROSITE: PS00280; BPTI\_KUNITZ\_1; 2.  
 CC PROSITE: PS00279; BPTI\_KUNITZ\_2; 2.  
 CC PROSITE: PS00213; LIPOCALIN; 1.  
 CC PROSITE: PS00213; LIPOCALIN; 1.  
 CC Glycoprotein; Plasma; Signal; Serine protease inhibitor; Repeat;  
 CC Lipocalin.  
 CC SIGNAL 1 19  
 CC CHAIN 20 203  
 CC CHAIN 206 352  
 CC  
 CC BINDING 53 53  
 CC DOMAIN 227 282  
 CC DOMAIN 283 348  
 CC DISULFID 91 188  
 CC DISULFID 231 281  
 CC DISULFID 240 264  
 CC DISULFID 256 277  
 CC DISULFID 287 337  
 CC DISULFID 296 320  
 CC DISULFID 312 333  
 CC DISULFID 241 343  
 CC ACT\_SITE 297 298  
 CC CARBOHYD 115 115  
 CC CARBOHYD 223 223  
 CC CARBOHYD 250 250  
 CC CARBOHYD 209 209  
 CC CONFLICT 217 217  
 CC CONFLICT 268 268  
 CC CONFLICT 274 274  
 CC CONFLICT 298 299  
 CC CONFLICT 330 330  
 CC CONFLICT 346 346  
 CC CARBOHYD 18 18  
 CC SEQUENCE 352 AA; 39235 MW; ED31C5CA02E70B19 CRC64;

Query Match 26.3%; Score 249.5; DB 1; Length 352;  
 Best Local Similarity 32.4%; Pred. No. 3.9e-16;  
 Matches 48; Conservative 16; Mismatches 45; Indels 39; Gaps 1;  
 9 DEFLYKRVGRKASMPHRYNTDSCGLFYVGGCDGNSNNLTRECLIKCAVTENA 68  
 229 DSCGLDYSGPGLCKRRRYNTSMACEFLVGGCGNGNPNLSKEKLCRTV---- 284  
 69 TGLIATSRNADSSVSPAPRQDSEHSDSMENVEEYCTANATGCRASFPRMYDVER 128  
 285 -----EACNLPVGGPCKRSTIQMLADPAVK 302  
 129 NSCNPFYGGCRGKNSYSEACMLRC 156  
 310 GKCVRFYGGKNGKNGFYSEKCEKVC 337  
 RESULT 2  
 AMBP\_MERUN STANDARD; PRT; 346 AA.  
 ID AMBP\_MERUN  
 AC Q62577; Q62576; 35, Created)  
 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 DT 15-JUL-1999 (Rel. 38, Last annotation update)  
 DE AMBP PROTEIN PRECURSOR (CONTAINS: ALPHA-1-MICROGLOBULIN; INTER-ALPHA-  
 DE TRYPSIN INHIBITOR LIGHT CHAIN (ITI-IC) (BIKUNIN) (HI-30)).  
 GN AMBP OR ITIL.  
 OS Meriones unguiculatus (Mongolian jird).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Gerbillinae;  
 OC Meriones.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE-LIVER:  
 RX MEDLINE-95110920; PubMed-7529051;  
 RA Ide H., Itoh H., Nawa Y.;  
 RT "Sequencing of cDNAs encoding alpha 1-microglobulin/bikunin of  
 RT Mongolian gerbil and Syrian golden hamster in comparison with man and  
 RT other species.";  
 RL Biochim. Biophys. Acta 1209:286-292(1994).  
 CC -1- FUNCTION: ALPHA-1-MICROGLOBULIN OCCURS IN MANY PHYSIOLOGICAL  
 CC FLUIDS INCLUDING PLASMA, URINE, AND CEREBROSPINAL FLUID. IT  
 CC APPEARS NOT ONLY AS A FREE MONOMER BUT ALSO IN COMPLEXES WITH IGA  
 CC AND ALBUMIN (BY SIMILARITY).  
 CC -1- FUNCTION: INTER-ALPHA-TRYPsin INHIBITOR, PRESENT IN PLASMA AND  
 CC URINE, INHIBITS TRYPSIN, PLASMIN, AND LYSOSOMAL GRANULOCYtic  
 CC ELASTASE (BY SIMILARITY).  
 CC -1- PTM: THE PRECURSOR IS PROTEOLYTICALLY PROCESSED INTO TWO  
 CC SEPARATELY FUNCTIONING PROTEINS.  
 CC -1- PTM: HC CONTAINS A COVALENTLY LINKED BROWN-YELLOW CHROMOPHORE (BY  
 CC SIMILARITY).  
 CC -1- SIMILARITY: IN THE N-TERMINAL SECTION, BELONGS TO THE LIPOCALIN  
 CC FAMILY.  
 CC -1- SIMILARITY: IN THE C-TERMINAL SECTION, BELONGS TO THE BPTI/KUNITZ  
 CC FAMILY OF INHIBITORS.  
 CC -----  
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 CC -----  
 CC EMBL: D31813; BAA06600.1; -  
 CC HSSP: P10646; IADZ.  
 CC INTERPRO: IPR000566; -  
 CC INTERPRO: IPR002223; -  
 CC INTERPRO: IPR002345; -  
 CC PFAM: PF00014; Kunitz\_BPTI; 2.  
 CC PFAM: PF00061; Lipocalin; 1.

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DR PRINTS: PR00179; LIPOCALIN.
DR PRINTS: PR00759; BASICPTASE.
DR PROSITE; PS00280; BPTI_KUNITZ_1; 2.
DR PROSITE; PS50279; BPTI_KUNITZ_2; 2.
KW Glycoprotein; Plasma; Signal; Serine protease inhibitor; Repeat;
LIPOCALIN.
FT SIGNAL 1 19
FT CHAIN 20 202
FT CHAIN 205 346
FT DOMAIN 226 281
FT DOMAIN 282 345
FT BINDING 52 52
FT DISULFID 90 187
FT DISULFID 230 280
FT DISULFID 239 263
FT DISULFID 255 276
FT DISULFID 286 336
FT DISULFID 295 319
FT DISULFID 311 332
FT CARBOHYD 114 114
FT CARBOHYD 249 249
FT ACT_SITE 240 241
FT ACT_SITE 296 297
FT ACT_SITE 346 AA; 38643 MW; F1A463810918D5F CRC64;
SQ SEQUENCE

Query Match
Best Local Similarity 32.4%; Score 247.5; DB 1; Length 346;
Matches 48; Conservative 15; Mismatches 46; Indels 39; Gaps 1;

QY 9 DFLVSKVYVGRGCRASMPRMWNTVTDSCQLFYGGCDGNSNNYLTKECLKCAVYENA 68
| : : : | : : : | : : : | : : : | : : : | : : : | : : : | : : : |
Db 228 DSCQLTYSRGPCLGMEKRIHNGTSMACEFYGGGLGNGNFISEKCLQTCRTVA-- 285
QY 69 TGDLATSRNAADSVSPAPRRDSEDSHSDMEYEEYCTANVTGFCRASFRWYDVDR 128
| : : : | : : : | : : : | : : : | : : : | : : : | : : : | : : : |
Db 286 -----CMLPYVGGFCRAVYIKLMDADAQ 308
QY 129 NSCNFFYGGCGRGNKNSYSEACMLRC 156
| : : : | : : : | : : : | : : : | : : : | : : : | : : : | : : : |
Db 309 GKCIQFTYGGCGKNGNKFSEKECKEYC 336

RESULT 3
FPI_RAT STANDARD: PRT; 302 AA.
AC 002445;
DT 01-JUL-1993 (Rel. 26, Created)
DT 01-JUL-1993 (Rel. 26, Last sequence update)
DT 30-MAY-2000 (Rel. 39, Last annotation update)
DE TISSUE FACTOR PATHWAY INHIBITOR PRECURSOR (TFPI) (LIPOPROTEIN-
DE ASSOCIATED COAGULATION INHIBITOR) (LACI) (EXTRINSIC PATHWAY INHIBITOR)
DE (EPI).
GN TFPI.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]
RC SEQUENCE FROM N.A.
RC STRAIN-SPRAGUE-DAWLEY; TISSUE-LIVER.
RX MEDLINE-92348361; PubMed-1639767;
RA Enjoji K.-I., Enli M., Mukai T., Kato H.;
RT "CDNA cloning and expression of rat tissue factor pathway inhibitor
RT (TFPI).";
RL J. Biochem. 111:681-687(1992).
-1- FUNCTION: INHIBITS FACTOR X (X(A)) DIRECTLY AND, IN A XA-DEPENDENT
WAY, INHIBITS VII(A)/TISSE FACTOR ACTIVITY, PREVIOUSLY BY FORMING
A QUATERNARY X(A)/LACI/VII(A)/TF COMPLEX. IT POSSESSES AN
ANTITHROMBOTIC ACTION AND ALSO THE ABILITY TO ASSOCIATE WITH
LIPOPROTEINS IN PLASMA.

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CC -1- TISSUE SPECIFICITY: MOST ABUNDANT IN HEART, LUNG, KIDNEY, AND
CC AORTIC ENDOTHELIAL CELLS.
CC -1- DOMAIN: THIS INHIBITOR CONTAINS THREE INHIBITORY DOMAINS.
CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.
CC HIGHLY SIMILAR TO TFPI2.
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-----
DR EMBL; D10926; BAA01724.1; -.
DR PIR; JX0213; TIRTKG.
DR HSSP; P10646; 1TFX.
DR INTERPRO; IPR002223; -.
DR PFAM; PF00014; Kunitz_BPTI; 3.
DR PRINTS; PR00759; BASICPTASE.
DR PROSITE; PS00280; BPTI_KUNITZ_1; 3.
DR PROSITE; PS50279; BPTI_KUNITZ_2; 3.
KW Serine protease inhibitor; Glycoprotein; Repeat; Blood coagulation;
KW Signal.
FT SIGNAL 1 28
FT CHAIN 29 302
FT CHAIN 53 103
FT DOMAIN 124 174
FT DOMAIN 222 272
FT DISULFID 53 103
FT DISULFID 62 86
FT DISULFID 78 99
FT ACT_SITE 63 64
FT ACT_SITE 124 174
FT DISULFID 133 157
FT DISULFID 149 170
FT ACT_SITE 134 135
FT DISULFID 222 272
FT DISULFID 231 255
FT DISULFID 247 268
FT ACT_SITE 232 233
FT CARBOHYD 144 144
FT CARBOHYD 251 251
FT CARBOHYD 261 261
SQ SEQUENCE 302 AA; 34554 MW; F9AE82130A24A59F CRC64;

Query Match
Best Local Similarity 34.0%; Score 246.5; DB 1; Length 302;
Matches 54; Conservative 21; Mismatches 65; Indels 19; Gaps 0;

QY 9 DFLVSKVYVGRGCRASMPRMWNTVTDSCQLFYGGCDGNSNNYLTKECLKCA-TYEN 67
| : : : | : : : | : : : | : : : | : : : | : : : | : : : | : : : |
Db 122 DFLVLEDPGICRGFMRYNNQSKQCEQFYGGCLGNSNFFLTDECRNTCDPVNEY 181
QY 68 ATGDLATSR-----NAADSVSPAPRRDSEDSHSDMEYEEYCTANVTGCGRA 117
| : : : | : : : | : : : | : : : | : : : | : : : | : : : | : : : |
Db 182 QKGDYVNTQITVDRITVNNVVIPOATKAPSDMDYDPS-----WCLEPADSGICKA 233
QY 118 SFRWYDVRRNSCNFFYGGCGRGNKNSYSEACMLRC 156
| : : : | : : : | : : : | : : : | : : : | : : : | : : : | : : : |
Db 234 SEKRFYTPAIGRCROFNYTGCGGNNNFTTKQDCNRAC 272

RESULT 4
IATR SHEEP STANDARD: PRT; 123 AA.
ID IATR SHEEP
AC P13371;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 01-APR-1990 (Rel. 14, Last annotation update)

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INTER-ALPHA-TRYPsin INHIBITOR (ITI) (GIR-14) (INHIBITORY FRAGMENT OF ITI) (FRAGMENT).

OS Ovis aries (Sheep), and Capra hircus (Goat).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovidae; Caprinae; Ovis.

[1]

RP SPECIES-SHEEP; Pubmed-2441725;

RC MEDLINE-87299012; Pubmed-2441725;

RA Rosp G., Hochstrasser K., Wachter E., Reisinger P.W.M.;

RT "The amino-acid sequence of the trypsin-released inhibitor from sheep inter-alpha-trypsin inhibitor.";

RL Biol. Chem. Hoppe-Seyler 368:727-731(1987).

[2]

RP SPECIES-HIRCUS; Pubmed-2481505;

RC MEDLINE-90105540; Pubmed-2481505;

RA Rosp G., Hochstrasser K., Gerl C., Wachter E.;

RT "Primary structure of a proteinase inhibitor released from goat serum inter-alpha-trypsin inhibitor.";

RL Biochim. Biophys. Acta 999:335-337(1989).

CC -1- FUNCTION: THIS INHIBITORY FRAGMENT, RELEASED FROM NATIVE ITI AFTER LIMITED PROTEOLYSIS WITH TRYPsin, CONTAINS TWO HOMOLOGOUS DOMAINS. WHEREAS THE SECOND DOMAIN IS A STRONG INHIBITOR OF TRYPsin, THE FIRST DOMAIN INTERACTS WEAKLY WITH PMN-GRANULOCYTIC ELASTASE AND NOT AT ALL WITH PANCREATIC ELASTASE.

CC -1- MISCELLANEOUS: THE AMINO ACID AT POSITION P2' (17) APPEARS TO DETERMINE THE SPECIFICITY OF THE INHIBITION OF DOMAIN 1. INHIBITORS WITH METHIONINE IN THIS POSITION INTERACT WEAKLY WITH CHYMOTRYPSIN AND ELASTASE; THOSE WITH LEUCINE INTERACT STRONGLY.

CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.

DR PIR. A29652; A29652.

DR HSSP; P10646; 1A02.

DR INTERPRO; IPR002223; -

DR PFAM; PF00014; Kunitz\_BPTI; 2.

DR PROSITE; PS00280; BPTI\_KUNITZ\_1; 2.

DR PROSITE; PS50279; BPTI\_KUNITZ\_2; 2.

KW Plasma: Glycoprotein; Serine protease inhibitor; Repeat.

FT NON-TER 1 56 I.

FT DOMAIN 57 123 II.

FT DISULFID 5 35 BY SIMILARITY.

FT DISULFID 14 38 BY SIMILARITY.

FT DISULFID 30 51 BY SIMILARITY.

FT DISULFID 61 111 BY SIMILARITY.

FT DISULFID 70 94 BY SIMILARITY.

FT DISULFID 86 107 BY SIMILARITY.

FT ACT\_SITE 15 16 INHIBITORY SITE (P1) (CHYMOTRYPSIN, ELASTASE).

FT ACT\_SITE 71 72 INHIBITORY SITE (P1) (TRYPsin).

FT CARBOHYD 24 24 N-LINKED (GLCNAc. . .).

FT NON-TER 123 123

SO SEQUENCE 123 AA; 13686 MW; 295038173F22D2D1 CRC64;

Query Match 25.88; Score 244.5; DB 1; Length 123;

Best Local Similarity 31.88; Pred. No. 3.5e-16;

Matches 47; Conservative 15; Mismatches 47; Indels 39; Gaps 1;

QY 9 DECIATSVKVRGRASMPRMWYVNTDGSQCLFYGGCDGNSNNYLTKECKLKCAITVENA 68

DB 3 DSCQLGYSGPCLGMFKRFRFYNGTSMACETFYGGCMGNNFPESEKELQTCRTV--- 58

QY 69 TGDLATSRMAADSVSPADPRDSDSHSDMNYEYCANAVTGPCRASFRWTEDYER 128

DB 59 -----OACNLPYRGKPCRGITELMAFDANK 83

QY 129 NSCNFTYGGCRGNKNSYSEACMLRC 156

DB 84 GRCVRFYGGCMGNGNOFTSQKCKRYC 111

RESULT 5

AMPB\_HUMAN STANDARD; PRT; 352 AA.

ID AMPB\_HUMAN P02759; P00977;

AC P02760; P02759; P00977;

DT 21-JUL-1986 (Rel. 01, Created)

DT 13-AUG-1987 (Rel. 05, Last sequence update)

DT 30-MAY-2000 (Rel. 39, Last annotation update)

DE AMPB PROTEIN PRECURSOR (CONTAINS: ALPHA-1-MICROGLOBULIN (PROTEIN HC) (COMPLEX-FORMING GLYCOPROTEIN HETEROGENEOUS IN CHARGE); INTER-ALPHA-TRYPsin INHIBITOR LIGHT CHAIN (ITI-LC) (BIKUNIN) (HI-30)).

DE AMPB OR ITI; OR HCP.

GN Homo sapiens (Human).

OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

[1]

RP SEQUENCE FROM N.A.

RC MEDLINE-91214554; Pubmed-1708673;

RA Vetr H., Gebhard W.;

RT "Structure of the human alpha 1-microglobulin-bikunin gene.";

RL Biol. Chem. Hoppe-Seyler 371:1185-1196(1990).

[2]

RP SEQUENCE FROM N.A.

RC TISSUE-LIVER; Pubmed-2430261;

RA MEDLINE-87040757; Pubmed-2430261;

RA Kaumeyer J.F., Polazzi J.O., Kotlick M.P.;

RT "The mRNA for a proteinase inhibitor related to the HI-30 domain of inter-alpha-trypsin inhibitor also encodes alpha-1-microglobulin (protein HC).";

RT Nucleic Acids Res. 14:7839-7850(1986).

[3]

RP SEQUENCE FROM N.A.

RC TISSUE-LIVER; Pubmed-1696200;

RA MEDLINE-90336621; Pubmed-1696200;

RA Diarra-Mehrpour M., Bourguignon J., Sesboue R., Salier J.P.;

RT "Structural analysis of the human inter-alpha-trypsin inhibitor light-chain gene.";

RT Eur. J. Biochem. 191:131-139(1990).

[4]

RP SEQUENCE OF 1-220 FROM N.A.

RC MEDLINE-86312901; Pubmed-2428011;

RA Traboni C., Cortese R.;

RT "Sequence of a full length cDNA coding for human protein HC (alpha 1 microglobulin).";

RL Nucleic Acids Res. 14:6340-6340(1986).

[5]

RP SEQUENCE OF 20-202 (INDIVIDUAL WITH TUBULAR PROTEINURIA).

RC MEDLINE-84126849; Pubmed-6198962;

RA Lopez C., Grubb A.O., Mendez E.;

RT "The complete amino acid sequence of human complex-forming glycoprotein heterogeneous in charge (protein HC) from one individual.";

RT Arch. Biochem. Biophys. 228:544-554(1984).

[6]

RP SEQUENCE OF 20-198 (VARIANT).

RC Lopez C., Grubb A.O., Mendez E.;

RT "Human protein HC displays variability in its carboxyl-terminal amino acid sequence.";

RL FEBS Lett. 144:349-353(1982).

[7]

RP SEQUENCE OF 20-198 (PATIENTS WITH TUBULAR PROTEINURIA).

RC MEDLINE-81184036; Pubmed-6164372;

RA Takagi T., Takagi K., Kawai T.;

RT "Complete amino acid sequence of human alpha 1-microglobulin.";

RL Biochem. Biophys. Res. Commun. 98:997-1001(1981).

[8]

RP SEQUENCE OF 206-350.

RC MEDLINE-85225968; Pubmed-2408638;

RA Reisinger P., Hochstrasser K., Albrecht G.J., Lempart K., Salier J.P.;

RT "Human inter-alpha-trypsin inhibitor: localization of the Kunitz-type domains in the N-terminal part of the molecule and their release as trypsin-like proteinase.";

RL Biol. Chem. Hoppe-Seyler 366:479-483(1985).

[9]  
 CARBOHYDRATE-LINKAGE SITES.  
 RX MEDLINE=82074265; PubMed=6171497;  
 RA Hochstrasser K., Schoenberger O.L., Rossmann I., Wachter E.;  
 RT "Kunitz-type proteinase inhibitors derived by limited proteolysis of  
 the inter-alpha-trypsin inhibitor, V. Attachments of carbohydrates in  
 the human urinary trypsin inhibitor isolated by affinity  
 chromatography.";  
 RL Hoppe-Seyler's Z. Physiol. Chem. 362:1357-1362(1981).  
 RN [10]  
 INHIBITORY SITE.  
 RX MEDLINE=85225940; PubMed=3890890;  
 RA Mori M., Travis J.;  
 RT "The reactive site of human inter-alpha-trypsin inhibitor is in the  
 amino-terminal half of the protein.";  
 RL Biol. Chem. Hoppe-Seyler 366:19-21(1985).  
 RN [11]  
 STRUCTURE OF CARBOHYDRATES.  
 MEDLINE=90306345; PubMed=1694784;  
 RA Escrichano J., Lopez-otin C., Hjerpe A., Grubb A., Mendez E.;  
 RT "Location and characterization of the three carbohydrate prosthetic  
 groups of human protein HC.";  
 RL FEBS Lett. 266:167-170(1990).  
 RN [12]  
 CHROMOPHORE.  
 TISSUE-URINE;  
 RX MEDLINE=91340714; PubMed=1714898;  
 RA Escrichano J., Grubb A., Calero M., Mendez E.;  
 RT "The protein HC chromophore is linked to the cysteine residue at  
 position 34 of the polypeptide chain by a reduction-resistant bond  
 and causes the charge heterogeneity of protein HC.";  
 RL J. Biol. Chem. 266:15758-15763(1991).  
 RN [13]  
 SEQUENCE OF 206-219, AND COVALENT LINKAGE WITH CHONDROITIN SULFATE.  
 TISSUE-PLASMA;  
 RX MEDLINE=94223087; PubMed=7513643;  
 RA Morille W., Capon C., Balduyck M., Sautiere P., Kouach M.,  
 Michalski C., Fournet B., Mizon J.;  
 RT "Chondroitin sulphate covalently cross-links the three polypeptide  
 chains of inter-alpha-trypsin inhibitor.";  
 RL Eur. J. Biochem. 221:881-888(1994).  
 RN [14]  
 SEQUENCE OF 206-223, AND CROSS-LINK SITE TO HC2.  
 MEDLINE=93232026; PubMed=7682553;  
 RA Engblid J.U., Salvesen G., Thøgersen I.B., Valnickova Z.,  
 Pizzo S.V., Hefta S.A.;  
 RT "Presence of the protein-glycosaminoglycan-protein covalent cross-link  
 in the inter-alpha-inhibitor-related proteinase inhibitor heavy chain  
 2/bikunin.";  
 RL J. Biol. Chem. 268:8711-8716(1993).  
 RN [15]  
 SEQUENCE OF 206-223, AND CROSS-LINK SITE TO HC3.  
 MEDLINE=91093267; PubMed=1898736;  
 RA Engblid J.U., Salvesen G., Hefta S.A., Thøgersen I.B.,  
 Rutherford S., Pizzo S.V.;  
 RT "Chondroitin 4-sulfate covalently cross-links the chains of the human  
 blood protein pre-alpha-inhibitor.";  
 RL J. Biol. Chem. 266:747-751(1991).  
 RN [16]  
 X-RAY CRYSTALLOGRAPHY (2.5 ANGSTROMS) OF 230-339.  
 MEDLINE=98227321; PubMed=9566199;  
 RA Xu Y., Carr P.D., Guss J.M., Ollis D.L.;  
 RT "The crystal structure of bikunin from the inter-alpha-inhibitor  
 complex: a serine protease inhibitor with two kunitz domains.";  
 RL J. Mol. Biol. 276:955-966(1998).  
 CC -1- FLUIDS INCLUDING PLASMA, URINE, AND CEREBROSPINAL FLUID IT  
 APPEARS NOT ONLY AS A FREE MONOMER BUT ALSO IN COMPLEXES WITH IGA  
 AND ALBUMIN.  
 CC -1- FUNCTION: INTER-ALPHA-TRYPSIN INHIBITOR, PRESENT IN PLASMA AND  
 URINE, INHIBITS TRYPSIN, PLASMIN, AND LISOSOMAL GRANULOCYTTIC  
 ELASTASE. ADDITIONAL PROTEOLYTIC PROCESSING IN THE KIDNEY AND/OR  
 URINE CAN PRODUCE FURTHER AMINO- AND CARBOXYL-END MODIFICATIONS

CC IN ITS SEQUENCE.  
 CC -1- SUBUNIT: INTER-ALPHA-TRYPSIN INHIBITOR CONSIST OF A LIGHT CHAIN  
 CC AND AN HEAVY CHAIN. THERE ARE THREE DIFFERENT HEAVY CHAINS.  
 CC -1- PTM: THE PRECURSOR IS PROTEOLYTICALLY PROCESSED INTO TWO  
 CC SEPARATELY FUNCTIONING PROTEINS.  
 CC -1- PTM: HC CONTAINS A COVALENTLY LINKED BROWN-YELLOW CHROMOPHORE.  
 CC -1- PTM: ADDITION OF GLYCOSAMINOGLYCAN CHONDROITIN SULFATE, ALONGS  
 CC MISCELLANEOUS: IN VITRO, THE FIRST TWELVE RESIDUES OF THE AMINO  
 CC END OF THE INHIBITOR APPEAR TO HAVE A REACTIVE SITE CAPABLE OF  
 CC INHIBITING THE ACTIVITY OF A NUMBER OF ENZYMES. ITS IN VIVO  
 CC FUNCTION IS NOT KNOWN.  
 CC -1- SIMILARITY: IN THE N-TERMINAL SECTION; BELONGS TO THE LIPOCALIN  
 CC FAMILY.  
 CC -1- SIMILARITY: IN THE C-TERMINAL SECTION; BELONGS TO THE BPTI/KUNITZ  
 CC FAMILY OF INHIBITORS.  
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 CC -----  
 DR EMBL: X54816; CAA38585.1; -  
 DR EMBL: X54817; CAA38585.1; JOINED.  
 DR EMBL: X54818; CAA38585.1; JOINED.  
 DR EMBL: X04225; CAA27803.1; -  
 DR EMBL: M88249; AAA59196.1; -  
 DR EMBL: M88165; AAA59196.1; JOINED.  
 DR EMBL: M88243; AAA59196.1; JOINED.  
 DR EMBL: M88244; AAA59196.1; JOINED.  
 DR EMBL: M88246; AAA59196.1; JOINED.  
 DR EMBL: M88247; AAA59196.1; JOINED.  
 DR EMBL: X04494; CAA28182.1; -  
 DR EMBL: X54817; CAA38586.1; -  
 DR PIR: A03217; HCHU  
 DR PIR: A25303; A25303.  
 DR PIR: S13433; S13433.  
 DR PIR: S10717; S10717.  
 DR PDB: 1B1K; 16-MAR-99.  
 DR SWISS-2DPAGE; P02760; HUMAN.  
 DR MIM: 176870; -  
 DR INTERPRO: IPR00566; -  
 DR INTERPRO: IPR002223; -  
 DR INTERPRO: IPR002345; -  
 DR PIR: PF00014; Kunitz-BPTI; 2.  
 DR PIR: PF00014; Kunitz-BPTI; 2.  
 DR PRINTS: PR00179; LIPOCALIN; 1.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 2.  
 DR PROSITE: PS02723; BPTI\_KUNITZ\_2; 2.  
 DR PROSITE: PS00213; LIPOCALIN; 1.  
 KW Glycoprotein; Plasma; Signal; Serine protease inhibitor; Repeat;  
 KW Lipocalin; 3D-structure.  
 Query Match 25.8%; Score 244.5; DB 1; Length 352;  
 Best Local Similarity 32.4%; Pred. No. 1,le-15;  
 Matches 48; Conservative 14; Mismatches 47; Indels 39; Gaps 1;  
 QY 9 DEFLVSVVGRCRASRMWYNTVDSQQLFVYGGCDGNSNNYLTKECKLKCAATVETNA 68  
 DB 229 DSQLGVSAGPCMGMTFRFYNGTSMACEFYFGCGAGNGNNGVTEKECLQCTRTVA-- 286  
 QY 69 TGLATSRNADSVSAPRROSESDHSDMFNYEYCTANAVTGPCRASEPRMYFVDER 128  
 DB 287 -----  
 QY 129 NSGNNTFYGGCRGNKNSYSSEACMLRC 156  
 DB 310 GRCVLFEPYGGCGGNGKRFYSSEKREYEC 337

RESULT 6  
TFPI\_RABIT STANDARD; PRT; 300 AA.  
ID TFPI\_RABIT  
AC P19761; Q28828;  
DT 01-FEB-1991 (Rel. 17, Created)  
DT 01-AUG-1992 (Rel. 23, Last sequence update)  
DT 01-NOV-1997 (Rel. 35, Last annotation update)  
DE TISSUE FACTOR PATHWAY INHIBITOR PRECURSOR (TFPI) (LIPOPROTEIN-  
DE ASSOCIATED COAGULATION INHIBITOR) (LACI) (EXTRINSIC PATHWAY INHIBITOR)  
DE (EPI).  
GN Oryctolagus cuniculus (Rabbit).  
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.  
RN  
RP  
RC TISSUE-LIVER.  
RC MEDLINE=91057146; PubMed=2136251;  
RA Wesselschmidt R.L., Girard T.O., Broze G.J. Jr.;  
"cDNA sequence of rabbit lipoprotein-associated coagulation  
inhibitor."  
Nucleic Acids Res. 18:6440-6440(1990).  
[2]  
REVIEWS TO 72; 211 AND 218.  
RN  
RP  
RC TISSUE-LIVER.  
RC MEDLINE=92335027; PubMed=1630940;  
RA Warr-Cramer B.J., Broze G.J. Jr., Komives E.A.;  
"cDNA sequence of rabbit tissue factor pathway inhibitor."  
Nucleic Acids Res. 20:3548-3548(1992).  
[3]  
SEQUENCE FROM N.A.  
RN  
RP  
RC TISSUE-LUNG.  
RC MEDLINE=93276427; PubMed=8503123;  
RA Belaouaj A., Kuppenswamy M.N., Birkhoff J.J., Bajaj S.P.;  
"Revised cDNA sequence of rabbit tissue factor pathway inhibitor."  
Thromb. Res. 69:547-553(1993).  
-1- FUNCTION: INHIBITS FACTOR X (X(A)) DIRECTLY AND, IN A XA-DEPENDENT  
MAY INHIBITS VII(A)/LACI/VIII(A)/TF COMPLEX. IT POSSESSES AN  
ANTITHROMBOTIC ACTION AND ALSO THE ABILITY TO ASSOCIATE WITH  
LIPOPROTEINS IN PLASMA. CONTAINS THREE INHIBITORY DOMAINS.  
-1- DOMAIN: THIS INHIBITOR CONTAINS THREE INHIBITORY DOMAINS.  
-1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
HIGHLY SIMILAR TO TFPI.  
-----  
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-----  
CC  
CC EMBL; X54708; CA38515.1; ALT-SEQ.  
CC EMBL; S61902; AAB26836.1; .  
CC  
CC  
CC EMBL; S61902; AAB26836.1; .  
CC  
CC  
CC HSSP; P10646; ITRX.  
CC INTERPRO; IPR002223; .  
CC PFM; PF00014; Kunitz\_BPTI; 3.  
CC PRINTS; PS00759; BASICPRASE.  
CC PROSITE; PS00280; BPTI\_KUNITZ\_1; 3.  
CC PROSITE; PS00279; BPTI\_KUNITZ\_2; 3.  
CC Serine protease inhibitor; Glycoprotein; Repeat; Blood coagulation;  
KW  
KW SIGNAL.  
FT  
FT CHAIN 1 24  
FT CHAIN 25 300  
FT CHAIN 50 100  
FT CHAIN 121 171  
FT CHAIN 213 263  
DOMAIN 121 171  
DOMAIN 213 263  
TISSUE FACTOR PATHWAY INHIBITOR.  
BPTI/KUNITZ INHIBITOR 1  
(VII(A)/TISSUE FACTOR BINDING SITE).  
BPTI/KUNITZ INHIBITOR 2  
(FACTOR X(A) BINDING SITE).  
BPTI/KUNITZ INHIBITOR 3  
(BY SIMILARITY).

FT DISULFID 50 100 BY SIMILARITY.  
FT DISULFID 59 83 BY SIMILARITY.  
FT DISULFID 75 96 REACTIVE BOND (BY SIMILARITY).  
FT ACT SITE 60 61 BY SIMILARITY.  
FT DISULFID 121 171 BY SIMILARITY.  
FT DISULFID 130 154 BY SIMILARITY.  
FT DISULFID 146 167 REACTIVE BOND (BY SIMILARITY).  
FT ACT SITE 131 132 BY SIMILARITY.  
FT DISULFID 213 263 BY SIMILARITY.  
FT DISULFID 222 246 BY SIMILARITY.  
FT DISULFID 238 259 REACTIVE BOND (BY SIMILARITY).  
FT ACT SITE 223 224 BY SIMILARITY.  
FT CARBOHYD 141 141 N-LINKED (GLCNAC. . .) (POTENTIAL).  
FT CARBOHYD 191 191 N-LINKED (GLCNAC. . .) (POTENTIAL).  
FT CARBOHYD 252 252 N-LINKED (GLCNAC. . .) (POTENTIAL).  
FT CARBOHYD 31 31 MISSING (IN REF. 3).  
FT CONFLICT 269 272 PKST -> RNL5 (IN REF. 3).  
FT CONFLICT 31 31 PKST -> RNL5 (IN REF. 3).  
FT SEQUENCE 300 AA; 34435 MM; A08DE36537708CA6 CRC64;

Query Match 25.7%; Score 244; DB 1; Length 300;  
Best Local Similarity 30.0%; Pred. No. 1,1e-15;  
Matches 48; Conservative 23; Mismatches 61; Indels 28; Gaps 2;

OY 4 ERSIHDFCLVSRVYGRCPASMPRMVNTDSCQLFYVGGDGSNNYLTRECLIKCAT 63  
DB 43 QKPTSFCAKMYVDGPCRAVYIKRFFENILTHQCEFFITGCGEGNREPSIECKCKCAR 102  
OY 64 VTENATGSLASRNADSVSPAPRRODESDHSSDMFYEEYCINAVTGCRASEFPKMY 123  
DB 103 DYFKMTTLITLTKYKGRPD-----FCFLDEDDGIORGIYIRNF 133  
OY 124 FVERNSCNFFTYGCGRKNKSYRSEACMLRCFROENP 163  
DB 139 YNNSKOCEREFYGGCIGLNNFESIECKNTC---ENP 174

RESULT 7  
AMBP\_PIG STANDARD; PRT; 337 AA.  
ID AMBP\_PIG  
AC P04366; P34954;  
DT 20-MAR-1987 (Rel. 04, Created)  
DT 01-FEB-1994 (Rel. 28, Last sequence update)  
DT 15-JUL-1999 (Rel. 38, Last annotation update)  
DE AMBP PROTEIN PRECURSOR (CONTAINS: ALPHA-1-MICROGLOBULIN; INTER-ALPHA-  
DE TRYPSIN INHIBITOR LIGHT CHAIN (ITI-LC) (HIKUNIN) (HI-30) (EI-14))  
DE (FRAGMENT).  
GN AMBP OR ITIL.  
OS Sus scrofa (Pig).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.  
RN  
RP  
RC MEDLINE=9035595; PubMed=1696914;  
RA Gebhard W., Schreimuehler T., Vetr H., Wachter E., Hochstrasser K.;  
"Complementary DNA and deduced amino acid sequences of porcine alpha  
1-microglobulin and bikunin."  
FEBS Lett. 269:32-36(1990).  
[2]  
RN  
RP  
RC TISSUE-LIVER.  
RC MEDLINE=91113729; PubMed=1703444;  
RA Tavares A.;  
"Molecular cloning of porcine alpha 1-microglobulin/HI-30 reveals  
developmental and tissue-specific expression of two variant messenger  
ribonucleic acids."  
RT Biochim. Biophys. Acta 1088:47-56(1991).  
[3]  
RN  
RP  
RC TISSUE-LIVER.  
RC MEDLINE=8525967; PubMed=2408637;  
RA Hochstrasser K., Wachter E., Albrecht G.J., Reisinger P.;  
"Kunitz-type proteinase inhibitors derived by limited proteolysis of  
the inter-alpha-trypsin inhibitor, X. The amino-acid sequences of the

RT trypsin-released inhibitors from horse and pig inter-alpha-trypsin  
 RL inhibitors.";  
 CC Biol. Chem. Hoppe-Seyler 366:473-478(1985).  
 CC -1- FUNCTION: ALPHA-1-MICROGLOBULIN OCCURS IN MANY PHYSIOLOGICAL  
 CC FLUIDS INCLUDING PLASMA, URINE, AND CEREBROSPINAL FLUID. IT  
 CC APPEARS NOT ONLY AS A FREE MONOMER BUT ALSO IN COMPLEXES WITH IGA  
 CC AND ALBUMIN.  
 CC -1- FUNCTION: INTER-ALPHA-TRYPsin INHIBITOR, PRESENT IN PLASMA AND  
 CC URINE, INHIBITS TRYPsin, PLASMIN, AND LYSOSOMAL GRANULOCYTIC  
 CC ELASTASE.  
 CC -1- P1M: THE PRECURSOR IS PROTEOLYTICALLY PROCESSED INTO TWO  
 CC SEPARATELY FUNCTIONING PROTEINS.  
 CC -1- P1M: HC CONTAINS A COVALENTLY LINKED BROWN-YELLOW CHROMOPHORE.  
 CC -1- SIMILARITY: IN THE N-TERMINAL SECTION; BELONGS TO THE LIPOCALIN  
 CC FAMILY.  
 CC -1- SIMILARITY: IN THE C-TERMINAL SECTION; BELONGS TO THE BPTI/KUNITZ  
 CC FAMILY OF INHIBITORS.  
 CC -----  
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 CC -----  
 CC EMBL: X53685; CAA37725.1; -;  
 CC EMBL: X52087; CAA36306.1; -;  
 CC DR PIR: A01208; TIPGR.  
 CC DR PIR: S11066; S11066.  
 CC DR HSSP: P10646; ITEX.  
 CC DR INTERPRO: IPR000566; -;  
 CC DR INTERPRO: IPR002223; -;  
 CC DR PFAM: PF00014; Kunitz\_BPTI; 2.  
 CC DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 2.  
 CC DR PROSITE: PS00279; BPTI\_KUNITZ\_2; 2.  
 CC DR PROSITE: PS00213; LIPOCALIN; 1.  
 CC KW Glycoprotein; Plasma; Signal; Serine protease inhibitor; Repeat;  
 CC LIPOCALIN.  
 CC FT NON\_TER 1 1  
 CC FT SIGNAL <1 4  
 CC FT CHAIN 5 188  
 CC FT CHAIN 191 337  
 CC FT  
 CC FT DOMAIN 212 267  
 CC FT BINDING 268 334  
 CC FT DISULFID 38 38  
 CC FT DISULFID 76 173  
 CC FT DISULFID 216 266  
 CC FT DISULFID 225 249  
 CC FT DISULFID 241 262  
 CC FT DISULFID 272 322  
 CC FT DISULFID 281 305  
 CC FT DISULFID 297 318  
 CC FT CARBOHYD 100 100  
 CC FT CARBOHYD 235 235  
 CC FT ACT\_SITE 226 227  
 CC FT  
 CC FT ACT\_SITE 282 283  
 CC FT CONFLICT 49 49  
 CC FT CONFLICT 259 259  
 CC FT CONFLICT 270 270  
 CC FT CONFLICT 278 278  
 CC FT CONFLICT 283 283  
 CC FT CONFLICT 285 286  
 CC FT CONFLICT 293 293  
 CC FT CONFLICT 311 311  
 CC FT CONFLICT 315 315  
 CC FT SEQUENCE 337 AA; 37690 MW; 1F630FE98E3CD70F CRC64;

Query Match

25.6%; Score 242.5; DB 1; Length 337;

Best Local Similarity 31.8%; Pred. No. 1,7e-15;  
 Matches 47; Conservative 19; Mismatches 43; Indels 39; Gaps 1;  
 QY 9 DCLVSKVYGRGRASPRMRYVVTGSCQLEFYGGCDGNSNNYLTRECLKRCATYENA 68  
 DB 214 DSCQGYSGPGLGMIKRFYNGSSMACETFFHYGCGMGNFVEKEDLCRYV---- 269  
 QY 69 TGDLATSRNADSVSPAPRRQSDHSDMNEYECIANAYTPCRASPRMYFVER 128  
 DB 270 -----EACSLPIVSGRCGFOLMADAVQ 294  
 QY 129 NSCNPFYGGCGKKNYSRSEACMLRC 156  
 DB 295 GKCVLENYGGQGNNGNOFYSEKECKEYC 322  
 RESULT 8  
 ID IATR\_HORSE STANDARD; PRT; 123 AA.  
 AC P04365;  
 DT 20-MAR-1987 (Rel. 04, Created)  
 DT 20-MAR-1987 (Rel. 04, Last sequence update)  
 DT 01-APR-1990 (Rel. 14, Last annotation update)  
 DE INTER-ALPHA-TRYPsin INHIBITOR (ITI) (HI-14) (INHIBITORY FRAGMENT OF  
 DE ITI) (FRAGMENT).  
 OS Equus caballus (Horse).  
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Perissodactyla; Equidae; Equus.  
 RN [1]  
 RP SEQUENCE.  
 RX MEDLINE-85225967; PubMed-2408637;  
 RA Hochstrasser K., Wachter E., Albrecht G.J., Reisinger P.;  
 RT "Kunitz-type proteinase inhibitors derived by limited proteolysis of  
 RT the inter-alpha-trypsin inhibitor, X. The amino-acid sequences of the  
 RT trypsin-released inhibitors from horse and pig inter-alpha-trypsin  
 RT inhibitors.";  
 RL Biol. Chem. Hoppe-Seyler 366:473-478(1985).  
 CC -1- FUNCTION: THIS INHIBITORY FRAGMENT, RELEASED FROM NATIVE ITI AFTER  
 CC LIMITED PROTEOLYSIS WITH TRYPsin, CONTAINS TWO HOMOLOGOUS DOMAINS.  
 CC WHEREAS THE SECOND DOMAIN IS A STRONG INHIBITOR OF TRYPsin, THE  
 CC FIRST DOMAIN INTERACTS WEAKLY WITH PMN-GRANULOCYTIC ELASTASE AND  
 CC NOT AT ALL WITH PANCREATIC ELASTASE.  
 CC -1- MISCELLANEOUS: THE AMINO ACID AT POSITION P2' (17) APPEARS TO  
 CC DETERMINE THE SPECIFICITY OF THE INHIBITION OF DOMAIN I.  
 CC INHIBITORS WITH METHIONINE IN THIS POSITION INTERACT WEAKLY WITH  
 CC CHYMOTRYPsin AND ELASTASE; THOSE WITH LEUCINE INTERACT STRONGLY.  
 CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 CC DR HSSP: P10646; IADZ.  
 CC DR INTERPRO: IPR002223; -;  
 CC DR PFAM: PF00014; Kunitz\_BPTI; 2.  
 CC DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 2.  
 CC DR PROSITE: PS00279; BPTI\_KUNITZ\_2; 2.  
 CC KW Plasma; Glycoprotein; Serine protease inhibitor; Repeat.  
 CC FT NON\_TER 1 1  
 CC FT DOMAIN 1 56  
 CC FT DOMAIN 57 123  
 CC FT DISULFID 5 55  
 CC FT DISULFID 14 38  
 CC FT DISULFID 30 51  
 CC FT DISULFID 61 111  
 CC FT DISULFID 70 94  
 CC FT DISULFID 86 107  
 CC FT ACT\_SITE 15 16  
 CC FT  
 CC FT ACT\_SITE 71 72  
 CC FT CARBOHYD 24 24  
 CC FT NON\_TER 123 123  
 CC FT SEQUENCE 123 AA; 13510 MW; CE1A9120774411D5 CRC64;

Query Match

25.5%; Score 241.5; DB 1; Length 123;  
 Best Local Similarity 31.8%; Pred. No. 6.6e-16;

DR	PfAM; PF00014; Kunitz-BPTI_2.
DR	PfAM; PF00061; Lipocalin_1.
DR	PRINTS; PR00179; LIPOCALIN.
DR	PRINTS; PRO0759; BASICPRASE.
DR	PROSITE; PS00280; BPTL_KUNITZ_1; 2.
DR	PROSITE; PSS0279; BPTL_KUNITZ_2; 2.
DR	PROSITE; PSS00213; LIPOCALIN_1.
KW	Glycoprotein; Plasma; Signal; Serine protease inhibitor; Repeat;
FT	Lipocalin..
FT	SIGNAL
FT	CHAIN
FT	CHAIN
FT	CHAIN
FT	DNAIN
FT	DNAIN
FT	BINDING
FT	DISULFD
FT	DISULFD
FT	DISULFD
FT	DISULFD
FT	DISULFD
FT	DISULFD
FT	CARBOHD
FT	CARBOND
FT	ACT_SITE
FT	ACT_SITE
FT	ACT_SITE
FT	CONFLICT
SO	SEQUENCE
FT	142
FT	142
FT	349 AA; 38851 MM; 1B7PB7DCB082A4E01 CRC64:
FT	BY SIMILARITY.
FT	ALPHA-1-MICROGLOBULIN.
FT	INTER-ALPHA-TRYPSIN INHIBITOR LIGHT CHAIN.
FT	I.
FT	II.
FT	CHROMOPHORE (BY SIMILARITY).
FT	BY SIMILARITY.
FT	BY SIMILARITY.
FT	BY SIMILARITY.
FT	BY SIMILARITY.
FT	BY SIMILARITY.
FT	BY SIMILARITY.
FT	N-LINKED (GLCNAC... ) (POTENTIAL).
FT	N-LINKED (GLCNAC... ) (POTENTIAL).
FT	INHIBITORY SITE (P1) (CHYMOTRYPsin,
FT	EASTASE) (BY SIMILARITY).
FT	INHIBITORY SITE (P1) (TRypsin) (BY
FT	SIMILARITY).
FT	G -> A (IN REF. 2).

RL J. Blochem. 115:708-714(1994).  
 CC -1- FUNCTION: INHIBITS FACTOR X (X(A)) DIRECTLY AND, IN A XA-DEPENDENT  
 CC WAY, INHIBITS VII(A)/TISSUE FACTOR ACTIVITY, PRESUMABLY BY FORMING  
 CC A QUATERNARY X(A)/LACI/VII(A)/TF COMPLEX. IT POSSESSES AN  
 CC ANTIHROMBOTIC ACTION AND ALSO THE ABILITY TO ASSOCIATE WITH  
 CC LIPOPROTEINS IN PLASMA.  
 CC -1- DOMAIN: THIS INHIBITOR CONTAINS THREE INHIBITORY DOMAINS.  
 CC -1- PTM: O-GLYCOSYLATED (BY SIMILARITY).  
 CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 CC HIGHLY SIMILAR TO TFP2.  
 CC -----  
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 CC -----  
 CC EMBL: S73337; AAB31955.1; -.  
 CC INTERPRO: IPR002223; -.  
 CC PFM: PFO0014; Kunitz\_BPTI; 3.  
 CC PRINTS: PR00759; BASICPTASE.  
 CC PROSITE: PS00280; BPTI\_KUNITZ\_1; 3.  
 CC PROSITE: PS00279; BPTI\_KUNITZ\_2; 3.  
 CC Serine protease inhibitor; Glycoprotein; Repeat; Blood coagulation;  
 CC Signal.  
 CC FT SIGNAL 1 28 BY SIMILARITY.  
 CC FT CHAIN 29 304 TISSUE FACTOR PATHWAY INHIBITOR.  
 CC FT DOMAIN 54 104 BPTI/KUNITZ INHIBITOR 1.  
 CC FT DOMAIN 125 175 (VII(A)/TISSUE FACTOR BINDING SITE).  
 CC FT DOMAIN 125 175 BPTI/KUNITZ INHIBITOR 2  
 CC (FACTOR X(A) BINDING SITE).  
 CC FT DOMAIN 217 267 (BPTI/KUNITZ INHIBITOR 3).  
 CC FT DISULFID 54 104 BY SIMILARITY.  
 CC FT DISULFID 63 87 BY SIMILARITY.  
 CC FT DISULFID 79 100 BY SIMILARITY.  
 CC FT ACT\_SITE 64 65 REACTIVE BOND (BY SIMILARITY).  
 CC FT DISULFID 125 175 BY SIMILARITY.  
 CC FT DISULFID 134 158 BY SIMILARITY.  
 CC FT DISULFID 150 171 BY SIMILARITY.  
 CC FT ACT\_SITE 135 136 REACTIVE BOND (BY SIMILARITY).  
 CC FT DISULFID 217 267 BY SIMILARITY.  
 CC FT DISULFID 226 250 BY SIMILARITY.  
 CC FT DISULFID 242 263 BY SIMILARITY.  
 CC FT ACT\_SITE 227 228 REACTIVE BOND (BY SIMILARITY).  
 CC FT CARBOHYD 145 145 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 CC FT CARBOHYD 195 195 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 CC FT CARBOHYD 256 256 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 CC SEQUENCE 304 AA: 35085 MW: 56133 kDa  
 CC -----

DT 01-NOV-1997 (Rel. 35, Created)  
 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 DT 15-JUL-1999 (Rel. 38, Last annotation update)  
 DE AMB PROTEIN PRECURSOR (CONTAINS: ALPHA-1-MICROGLOBULIN; INTER-ALPHA-  
 DE TRYPsin INHIBITOR LIGHT CHAIN (ITI-IC) (BIKUNIN) (HI-30)).  
 GN AMB OR ITIL.  
 OS Mesocricetus auratus (Golden hamster).  
 CC Eukaryota; Metazoa; Chordata; Craniata; Euteleostomi;  
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Cricetidae;  
 CC Mesocricetus.  
 CC [1]  
 RN SEQUENCE FROM N.A.  
 RP TISSUE=LIVER;  
 RX MEDLINE=95110820; PubMed=7529051;  
 RA Ide H., Itoh H., Nawa Y.;  
 RT "Sequencing of cDNAs encoding alpha 1-microglobulin/bikunin of  
 RT Mongolian gerbil and Syrian golden hamster in comparison with man and  
 RT other species.";  
 RL Blochem. Biophys. Acta 1209:286-292(1994).  
 CC -1- FUNCTION: ALPHA-1-MICROGLOBULIN OCCURS IN MANY PHYSIOLOGICAL  
 CC FLUIDS INCLUDING PLASMA, URINE, AND CEREBROSPINAL FLUID. IT  
 CC APPEARS NOT ONLY AS A FREE MONOMER BUT ALSO IN COMPLEXES WITH TGA  
 CC AND ALBUMIN (BY SIMILARITY).  
 CC -1- FUNCTION: INTER-ALPHA-TRYPsin INHIBITOR, PRESENT IN PLASMA AND  
 CC URINE, INHIBITS TRYPsin, PLASMIN, AND LYSOSOMAL GRANULOCYTIC  
 CC ELASTASE (BY SIMILARITY).  
 CC -1- PTM: THE PRECURSOR IS PROTEOLYTICALLY PROCESSED INTO TWO  
 CC SEPARATELY FUNCTIONING PROTEINS.  
 CC -1- PTM: HC CONTAINS A COVALENTLY LINKED BROWN-YELLOW CHROMOPHORE (BY  
 CC SIMILARITY).  
 CC -1- SIMILARITY: IN THE N-TERMINAL SECTION, BELONGS TO THE LIPOCALIN  
 CC FAMILY.  
 CC -1- SIMILARITY: IN THE C-TERMINAL SECTION, BELONGS TO THE BPTI/KUNITZ  
 CC FAMILY OF INHIBITORS.  
 CC -----  
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 CC -----  
 CC EMBL: D31814; BAA06601.1; -.  
 CC HSSP: P10646; 1TFX.  
 CC DR INTERPRO: IPR000566; -.  
 CC DR INTERPRO: IPR002223; -.  
 CC DR INTERPRO: IPR002345; -.  
 CC DR PFM: PFO0014; Kunitz\_BPTI; 2.  
 CC DR PFM: PFO0061; Lipocalin; 1.  
 CC DR PRINTS: PR00179; Lipocalin.  
 CC DR PRINTS: PR00759; BASICPTASE.  
 CC DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 2.  
 CC DR PROSITE: PS00279; BPTI\_KUNITZ\_2; 2.  
 CC DR PROSITE: PS00213; Lipocalin; 1.  
 CC KW Glycoprotein; Plasma; Signal; Serine protease inhibitor; Repeat;  
 CC Lipocalin.  
 CC FT SIGNAL 1 19 BY SIMILARITY.  
 CC FT CHAIN 20 202 ALPHA-1-MICROGLOBULIN.  
 CC FT CHAIN 205 349 INTER-ALPHA-TRYPsin INHIBITOR LIGHT  
 CC CHAIN.  
 CC FT DOMAIN 226 281 I.  
 CC FT DOMAIN 282 348 II.  
 CC FT BINDING 52 52 CHROMOPHORE (BY SIMILARITY).  
 CC FT DISULFID 90 187 BY SIMILARITY.  
 CC FT DISULFID 230 280 BY SIMILARITY.  
 CC FT DISULFID 239 263 BY SIMILARITY.  
 CC FT DISULFID 255 276 BY SIMILARITY.  
 CC FT DISULFID 286 336 BY SIMILARITY.  
 CC FT DISULFID 295 319 BY SIMILARITY.  
 CC FT DISULFID 311 332 BY SIMILARITY.  
 CC FT CARBOHYD 35 35 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 CC FT CARBOHYD 114 114 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 CC -----

FT ACT\_SITE 240 241 INHIBITORY SITE (P1) (CHYMOTRYPSIN,  
 FT ACT\_SITE 296 297 ELASTASE) (BY SIMILARITY).  
 FT ACT\_SITE 296 297 INHIBITORY SITE (P1) (TRYPSIN) (BY  
 FT SEQUENCE 349 AA; 38782 MW; 8C954584B7DBE28 CRC64;  
 SO  
 Query Match 24.8%; Score 235.5; DB 1; Length 349;  
 Best Local Similarity 31.1%; Pred. No. 7.8e-15;  
 Matches 46; Conservative 17; Mismatches 46; Indels 39; Gaps 1;

QY 9 DECLVSKVYGRGRASMPRMWYNTDSCOLFVYGGDGNMNYLTKRECKKCATVTEA 68  
 DB 228 DSCQLYSSEGPLGMEKRYTYNGASMACETFFHYGGCLGNMNFNSEKCIQTCTVAA-- 285  
 QY 69 TGDLATSRNAADSVSPAPRRDSEDDHSDMFYEEYCTANAVTGCRASFPMWYDVER 128  
 DB 286 -----CSLPYVQGPCRAVYELMAFDAAQ 308

129 NSCNNTFYGGCGKNGKNSYSEACMLRC 156  
 309 GKCVDFSYGGCKGNGKMFYSEKCEKCYC 336

RESULT 12  
 ABB\_MOUSE STANDARD: PRT; 349 AA.  
 ID ABB\_MOUSE  
 AC 007456; 061294;  
 DT 01-NOV-1997 (Rel. 35, Created)  
 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 DT 15-JUL-1999 (Rel. 38, Last annotation update)  
 DE ABB PROTEIN PRECURSOR [CONTAINS: ALPHA-1-MICROGLOBULIN; INTER-ALPHA-  
 DE TRYPSIN INHIBITOR LIGHT CHAIN (ITI-LC) (BIKUNIN) (HI-30)].  
 GN ABB OR ITIL.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 RN  
 RN [1]  
 RC SEQUENCE FROM N.A.  
 RC STRAIN-BALB/C; TISSUE-LIVER;  
 RX MEDLINE-93363639; PubMed-7689339;  
 RA Chan P., Salier J.P.;  
 RT "Mouse alpha-1-microglobulin/Bikunin precursor: CDNA analysis, gene  
 RT evolution and physical assignment of the gene next to the orosomucoid  
 RT locus.";  
 RL Biochim. Biophys. Acta 1174:195-200(1993).  
 RN  
 RN [2]  
 RC SEQUENCE FROM N.A.  
 RC STRAIN-C57BL/6; TISSUE-LIVER;  
 RX MEDLINE-95189774; PubMed-7533761;  
 RA Itoh H., Ide H., Kataoka H., Tomita M., Yoshihara H., Nawa Y.;  
 RT "CDNA sequencing of mouse alpha 1-microglobulin/inter-alpha-trypsin  
 RT inhibitor light chain and its expression in acute inflammation.";  
 RL J. Biochem. 116:767-772(1994).  
 RN  
 RN [3]  
 RC SEQUENCE OF 128-349 FROM N.A.  
 RC STRAIN-C57BL/6; TISSUE-LIVER;  
 RA Itoh H., Ide H., Yoshihara H., Nawa Y.;  
 RT Submitted (JAN-1994) to the EMBL/Genbank/DBJ databases.  
 CC -1- FUNCTION: ALPHA-1-MICROGLOBULIN OCCURS IN MANY PHYSIOLOGICAL  
 CC FLUIDS INCLUDING PLASMA, URINE, AND CEREBROSPINAL FLUID. IT  
 CC APPEARS NOT ONLY AS A FREE MONOMER BUT ALSO IN COMPLEXES WITH IGA  
 CC AND ALBUMIN (BY SIMILARITY).  
 CC -1- FUNCTION: INTER-ALPHA-TRYPSIN INHIBITOR, PRESENT IN PLASMA AND  
 CC URINE, INHIBITS TRYPSIN, PLASMIN, AND LYSOSOMAL GRANULOCYTIC  
 CC ELASTASE (BY SIMILARITY).  
 CC -1- SUBUNIT: INTER-ALPHA-TRYPSIN INHIBITOR CONSIST OF A LIGHT CHAIN  
 CC AND AN HEAVY CHAIN. THERE ARE THREE DIFFERENT HEAVY CHAINS.  
 CC -1- PTM: THE PRECURSOR IS PROTEOLYTICALLY PROCESSED INTO TWO  
 CC SEPARATELY FUNCTIONING PROTEINS.  
 CC -1- PTM: HC CONTAINS A COVALENTLY LINKED BROWN-YELLOW CHROMOPHORE (BY  
 CC SIMILARITY).  
 CC -1- SIMILARITY: IN THE N-TERMINAL SECTION; BELONGS TO THE LIPOCALIN

CC FAMILY.  
 CC -1- SIMILARITY: IN THE C-TERMINAL SECTION; BELONGS TO THE BPTI/KUNITZ  
 CC FAMILY OF INHIBITORS.  
 CC  
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 CC or send an email to [license@isb.slb.ch](mailto:license@isb.slb.ch)).  
 CC  
 CC EMBL: X68680; CAA48640.1; -;  
 CC DR D28812; BAA05973.1; -;  
 CC DR HSP: P12111; 1KUN.  
 CC DR MGD: MGI:88002; ABBP.  
 CC DR INTERPRO: IPR000566; -;  
 CC DR INTERPRO: IPR002223; -;  
 CC DR INTERPRO: IPR002345; -;  
 CC DR PRAM: PF00014; Kunitz\_BPTI; 2.  
 CC DR PRAM: PF00061; Lipocalin; 1.  
 CC DR PRINTS: PR00179; LIPOCALIN.  
 CC DR PRINTS: PR00759; BASICPTASE.  
 CC DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 2.  
 CC DR PROSITE: PS00279; BPTI\_KUNITZ\_2; 2.  
 CC DR PROSITE: PS00213; LIPOCALIN; 1.  
 CC KW Glycoprotein; Plasma; Signal; Serine protease inhibitor; Repeat;  
 KW Lipocalin.  
 FT SIGNAL 1 19 BY SIMILARITY.  
 FT CHAIN 20 202 ALPHA-1-MICROGLOBULIN.  
 FT CHAIN 205 349 INTER-ALPHA-TRYPSIN INHIBITOR LIGHT  
 FT CHAIN  
 FT DOMAIN 226 281 I.  
 FT BINDING 282 348 II.  
 FT BINDING 52 52 CHROMOPHORE (BY SIMILARITY).  
 FT DISULFID 90 187 BY SIMILARITY.  
 FT DISULFID 230 280 BY SIMILARITY.  
 FT DISULFID 239 263 BY SIMILARITY.  
 FT DISULFID 255 276 BY SIMILARITY.  
 FT DISULFID 286 336 BY SIMILARITY.  
 FT DISULFID 295 319 BY SIMILARITY.  
 FT DISULFID 311 332 BY SIMILARITY.  
 FT CARBOHYD 33 33 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT CARBOHYD 114 114 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT CARBOHYD 233 233 INHIBITORY SITE (P1) (CHYMOTRYPSIN,  
 FT ACT\_SITE 240 241 ELASTASE) (BY SIMILARITY).  
 FT ACT\_SITE 296 297 INHIBITORY SITE (P1) (TRYPSIN) (BY  
 FT ACT\_SITE  
 FT CONFLICT 65 65 Q -> S (IN REF. 2).  
 FT SEQUENCE 349 AA; 39070 MW; CE4D9C7375DA80B CRC64;  
 SQ

Query Match 24.8%; Score 235.5; DB 1; Length 349;  
 Best Local Similarity 29.7%; Pred. No. 7.8e-15;  
 Matches 44; Conservative 18; Mismatches 47; Indels 39; Gaps 1;

QY 9 DECLVSKVYGRGRASMPRMWYNTDSCOLFVYGGDGNMNYLTKRECKKCATVTEA 68  
 DB 228 DSCQLYSSEGPLGMEKRYTYNGASMACETFFHYGGCLGNMNFNSEKCIQTCTVAA-- 285  
 QY 69 TGDLATSRNAADSVSPAPRRDSEDDHSDMFYEEYCTANAVTGCRASFPMWYDVER 128  
 DB 286 -----CSLPYVQGPCRAVYELMAFDAAQ 308

129 NSCNNTFYGGCGKNGKNSYSEACMLRC 156  
 309 GKCVDFSYGGCKGNGKMFYSEKCEKCYC 336

RESULT 13  
 TFPI\_HUMAN STANDARD: PRT; 304 AA.  
 ID TFPI\_HUMAN



AC p10646;  
 DT 01-JUL-1989 (Rel. 11, Created)  
 DT 01-JUL-1989 (Rel. 11, Last sequence update)  
 DT 15-DEC-1998 (Rel. 37, Last annotation update)  
 DE TISSUE FACTOR PATHWAY INHIBITOR PRECURSOR (TFPI) (LIPOPROTEIN-  
 DE ASSOCIATED COAGULATION INHIBITOR) (LACI) (EXTRINSIC PATHWAY INHIBITOR)  
 GN TFPI OR TFPI OR LACI.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 RP [1]  
 RP MEDLINE=88198127; PubMed=2452157;  
 RA Mun T.-C., Kretzmer K.K., Girard T.J., Miletich J.P., Broze G.J. Jr.;  
 RT Cloning and characterization of a cDNA coding for the lipoprotein-  
 RT associated coagulation inhibitor of a CDNA coding for the lipoprotein-  
 RT tandem Kunitz-type inhibitory domains.";  
 RT J. Biol. Chem. 263:6001-6004(1988).  
 RL [2]  
 RL MEDLINE=91129227; PubMed=1993173;  
 RA van der Logt C.P.E., Reitsma P.H., Bertina R.M.;  
 RT "Intron-exon organization of the human gene coding for the  
 RT lipoprotein-associated coagulation inhibitor: the factor xa dependent  
 RT inhibitor of the extrinsic pathway of coagulation.";  
 RT Biochemistry 30:1571-1577(1991).  
 RN [3]  
 RN MEDLINE=91161593; PubMed=2002045;  
 RA Girard T.J., Eddy R., Wesselschmidt R.L., Macphail L.A.,  
 RT Likert K.M., Byers M.G., Shows T.B., Broze G.J. Jr.;  
 RT "Structure of the human lipoprotein-associated coagulation inhibitor  
 RT gene. Intron/exon gene organization and localization of the gene to  
 RT chromosome 2.";  
 RT J. Biol. Chem. 266:5036-5041(1991).  
 RN [4]  
 RN MEDLINE=89388722; PubMed=2781520;  
 RA Girard T.J., Warren L.A., Novotny W.F., Broze G.J. Jr.;  
 RT "Identification of the 1.4 kb and 4.0 kb messages for the lipoprotein  
 RT associated coagulation inhibitor and expression of the encoded  
 RT protein.";  
 RT Thromb. Res. 55:37-50(1989).  
 RN [5]  
 RN MEDLINE=90036966; PubMed=2553722;  
 RA Novotny W.F., Girard T.J., Miletich J.P., Broze G.J. Jr.;  
 RT "Purification and characterization of the lipoprotein-associated  
 RT coagulation inhibitor from human plasma.";  
 RT J. Biol. Chem. 264:18832-18837(1989).  
 RN [6]  
 RN MEDLINE=89181950; PubMed=2927510;  
 RA Girard T.J., Warren L.A., Novotny W.F., Likert K.M., Brown S.G.,  
 RT Miletich J.P., Broze G.J. Jr.;  
 RT "Functional significance of the Kunitz-type inhibitory domains of  
 RT lipoprotein-associated coagulation inhibitor.";  
 RT Nature 338:518-520(1989).  
 RN [7]  
 RN MEDLINE=96224851; PubMed=8639592;  
 RA Nakahara Y., Miyata T., Hamuro T., Funatsu A., Miyagi M.,  
 RT "Amino acid sequence and carbohydrate structure of a recombinant  
 RT human tissue factor pathway inhibitor expressed in Chinese hamster  
 RT ovary cells: one N- and two O-linked carbohydrate chains are located  
 RT between Kunitz domains 2 and 3 and one N-linked carbohydrate chain is  
 RT in Kunitz domain 2.";  
 RL Biochemistry 35:6450-6459(1996).  
 RN [8]  
 RN REVIEW.

RX MEDLINE=91104709; PubMed=2271516;  
 RA Broze G.J. Jr., Girard T.J., Novotny W.F.;  
 RT "Regulation of coagulation by a multivalent Kunitz-type inhibitor.";  
 RL Biochemistry 29:7539-7546(1990).  
 RN [9]  
 RN STRUCTURE BY NMR OF 121-182.  
 RX MEDLINE=97342711; PubMed=9199408;  
 RA Burgerling H.J., Ordonez L.P., van der Doelen A., Mulders J.,  
 RT Theunissen H.J., Grootehuis P.D., Bode W., Huber R., Stubbs M.T.;  
 RT "The second Kunitz domain of human tissue factor pathway inhibitor:  
 RT cloning, structure determination and interaction with factor Xa.";  
 RL J. Mol. Biol. 269:395-407(1997).  
 CC -1- FUNCTION: INHIBITS FACTOR X (X(A)) DIRECTLY AND, IN A XA-DEPENDENT  
 CC WAY, INHIBITS VII(A)/TISSUE FACTOR ACTIVITY, PRESUMABLY BY FORMING  
 CC A QUATERNARY X(A)/LACI/VII(A)/TF COMPLEX. IT POSSESSES AN  
 CC ANTIHEMOSTATIC ACTION AND ALSO THE ABILITY TO ASSOCIATE WITH  
 CC LIPOPROTEINS IN PLASMA.  
 CC -1- TISSUE SPECIFICITY: MOSTLY IN ENDOTHELIAL CELLS.  
 CC -1- DOMAIN: THIS INHIBITOR CONTAINS THREE INHIBITORY DOMAINS.  
 CC -1- PTM: O-GLYCOSYLATED.  
 CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 CC HIGHLY SIMILAR TO TFPI2.  
 CC -----  
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 CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).  
 CC -----  
 CC EMBL: J03225; AAA52022.1; -  
 CC EMBL: M58650; AAA59480.1; -  
 CC EMBL: M58644; AAA59480.1; JOINED.  
 CC EMBL: M58645; AAA59480.1; JOINED.  
 CC EMBL: M58646; AAA59480.1; JOINED.  
 CC EMBL: M58647; AAA59480.1; JOINED.  
 CC EMBL: M58648; AAA59480.1; JOINED.  
 CC EMBL: M58649; AAA59480.1; JOINED.  
 CC EMBL: M59493; AAA59526.1; JOINED.  
 CC EMBL: M59494; AAA59526.1; JOINED.  
 CC EMBL: M59495; AAA59526.1; JOINED.  
 CC EMBL: M59496; AAA59526.1; JOINED.  
 CC EMBL: M59497; AAA59526.1; JOINED.  
 CC EMBL: M59498; AAA59526.1; JOINED.  
 CC PTR: A28650; TIRUGK.  
 CC PTR: A34315; A34315.  
 CC PTR: A60433; A60433.  
 CC PTR: S03903; S03903.  
 CC PDB: 1ADZ; 25-FEB-98.  
 CC PDB: 1TFX; 21-JAN-98.  
 CC MIM: 152310; -  
 CC INTERPRO: IPR002223; -  
 CC PRAM: PF00014; Kunitz\_BPTI; 3.  
 CC PRINTS: PR00759; BASICPTASE.  
 CC PROSITE: PS00289; BPTI\_KUNITZ\_1; 3.  
 CC PROSITE: PS00279; BPTI\_KUNITZ\_2; 3.  
 CC KW Serine protease inhibitor; Glycoprotein; Repeat; Blood coagulation;  
 CC SIGNAL; 3D-structure.  
 CC FT CHAIN 1 28  
 CC FT 29 304  
 CC FT DOMAIN 54 104  
 CC FT  
 CC DOMAIN 125 175  
 CC FT  
 CC DOMAIN 217 267  
 CC FT DISULFID 54 104  
 CC FT DISULFID 63 87  
 CC FT DISULFID 79 100  
 CC FT ACT\_SITE 64 65  
 CC FT DISULFID 125 175  
 CC FT DISULFID 134 158  
 CC  
 CC TISSUE FACTOR PATHWAY INHIBITOR.  
 CC BPTI/KUNITZ INHIBITOR 1  
 CC (VII(A)/TISSUE FACTOR BINDING SITE).  
 CC BPTI/KUNITZ INHIBITOR 2  
 CC (FACTOR X(A) BINDING SITE).  
 CC BPTI/KUNITZ INHIBITOR 3.  
 CC BY SIMILARITY.  
 CC BY SIMILARITY.  
 CC BY SIMILARITY.  
 CC REACTIVE BOND (BY SIMILARITY).  
 CC BY SIMILARITY.  
 CC BY SIMILARITY.

FT DISULFID 150 177 BY SIMILARITY.  
 FT ACT\_SITE 135 136 REACTIVE BOND (BY SIMILARITY).  
 FT DISULFID 217 267 BY SIMILARITY.  
 FT DISULFID 226 250 BY SIMILARITY.  
 FT DISULFID 242 263 BY SIMILARITY.  
 FT ACT\_SITE 227 228 REACTIVE BOND (BY SIMILARITY).  
 FT CARBOHYD 145 145 N-LINKED (GLCNAC. . .).  
 FT CARBOHYD 202 202  
 FT CARBOHYD 203 203  
 FT CARBOHYD 195 195 N-LINKED (GLCNAC. . .).  
 FT CARBOHYD 64 64 K->I: ABOLISHES INHIBITION OF VII(A)/TF.  
 FT MUTAGEN 135 135 R->L: ABOLISHES INHIBITION OF X(A).  
 FT MUTAGEN 227 227 R->L: ABOLISHES INHIBITION OF VII(A)/TF.  
 FT SEQUENCE 304 AA: 35015 MW: 5281E32B758B44FE CRC64;  
 SO

Query Match 24.6%; Score 233.5; DB 1; Length 304;  
 Best Local Similarity 33.6%; Pred. No. 1e-14; Indels 11; Gaps 3;  
 Matches 51; Conservative 26; Mismatches 64

9 DCLVSKVYVGRASMPRMWYNTDSCGLFYGGCGDGSNNNYLTKKCKATVTEA 68  
 123 DCFLEEDGICRGYTRFYNNOTKOCERFKYGGGLGMMNFTLECKNIC---EDGP 179  
 69 TG----DLATSRNAADSVSPAPRRDSEDHSDMFNYEYCTANAYTGPCRASFRMYE 124  
 180 NGQVQNYGQGLNANVNSLTP---QSTKPSLFEFFHGSWCLTPADRGICRANENRFY 235  
 125 DVERNSCNFTYGGCGKGNKSYSEECMLRC 156  
 236 NSVIGKCRPFKYSKCGGNNENFTSKOCLNAC 267

RESULT 14  
 ID TP2\_HUMAN STANDARD; PRT: 235 AA.  
 AC P48307;  
 DT 01-FEB-1996 (Rel. 33, Created)  
 DT 01-FEB-1996 (Rel. 33, Last sequence update)  
 DT 01-NOV-1997 (Rel. 35, Last annotation update)  
 DE TISSUE FACTOR PATHWAY INHIBITOR 2 PRECURSOR (TFPI-2) (PLACENTAL  
 DE PROTEIN 5) (PP5).  
 GN TFPI2.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 RN [1]  
 RP SEQUENCE FROM N.A. AND PARTIAL SEQUENCE.  
 RA TISSUE-PLACENTA; PubMed-7896752;  
 RA MEDLINE-95204397; PubMed-7896752;  
 RA Miyagi Y., Koshikawa N., Yasumitsu H., Miyagi E., Hirahara F.,  
 Aoki I., Misugi K., Umeda M., Miyazaki K.,  
 "cDNA cloning and mRNA expression of a serine proteinase inhibitor  
 secreted by cancer cells: identification as placental protein 5 and  
 tissue factor pathway inhibitor-2.";  
 RT J. Biochem. 116:939-942(1994).  
 RL J.  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RP TISSUE-PLACENTA;  
 RX MEDLINE-94211862; PubMed-8159751;  
 RA Sprechter C.A., Kistiel W., Mathewes S., Foster D.C.;  
 "Molecular cloning, expression, and partial characterization of a  
 second human tissue-factor-pathway inhibitor.2.";  
 RT Proc. Natl. Acad. Sci. U.S.A. 91:3353-3357(1994).  
 RL [3]  
 RN SEQUENCE FROM N.A.  
 RP Magi L.;  
 RA Submitted (MAY-1997) to the EMBL/Genbank/DBJ databases.  
 RP PARTIAL SEQUENCE OF 23-35; 47-53 AND 133-146.  
 RP TISSUE-PLACENTA;  
 RX MEDLINE-8810628; PubMed-3276312;  
 RA Buetzow R., Huhtala M.-L., Bohn H., Virtanen I., Seppaelae M.;

RT "Purification and characterization of placental protein 5.";  
 RL Biochem. Biophys. Res. Commun. 150:483-490(1988).  
 RN [5]  
 RP ERRATUM.  
 RA Buetzow R., Huhtala M.-L., Bohn H., Virtanen I., Seppaelae M.;  
 Biochem. Biophys. Res. Commun. 151:630-631(1988).  
 RL Biochem. Biophys. Res. Commun. 151:630-631(1988).  
 CC -1- FUNCTION: SEEMS TO INHIBIT TRYPSIN, FACTOR VII(A)/TISSUE FACTOR,  
 CC WEAKLY FACTOR XA. HAS NO EFFECT ON THROMBIN.  
 CC -1- TISSUE SPECIFICITY: UMBILICAL VEIN ENDOTHELIAL CELLS, LIVER,  
 CC PLACENTA, HEART, PANCREAS, AND MATERNAL SERUM AT ADVANCED  
 CC PREGNANCY.  
 CC -1- DOMAIN: THIS INHIBITOR CONTAINS THREE INHIBITORY DOMAINS.  
 CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 CC HIGHLY SIMILAR TO TPFI.  
 CC -----  
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 CC or send an email to [license@sib.ch](mailto:license@sib.ch)).  
 CC -----  
 CC DR EMBL: D29992; BA06272.1; .  
 CC DR EMBL: L27624; AA20094.1; .  
 CC DR EMBL: AC002076; AAB54049.1; .  
 CC DR PIR: A34029; A34029.  
 CC DR PIR: B34029; B34029.  
 CC DR PIR: C34029; C34029.  
 CC DR HSP: P12111; IKNY.  
 CC DR MIM: 600033; .  
 CC DR INTERPRO: IPR002223; .  
 CC DR PFAM: PF00014; Kunitz\_BPTI; 3.  
 CC DR PRINTS: PR00759; BASICTPASE.  
 CC DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 2.  
 CC DR PROSITE: PS00279; BPTI\_KUNITZ\_2; 3.  
 CC DR PROSITE: PS00279; BPTI\_KUNITZ\_2; 3.  
 CC KW Serine protease inhibitor; Glycoprotein; Repeat; signal;  
 CC KW Blood coagulation.  
 CC FT SIGNAL 1 22  
 CC FT CHAIN 23 25 TISSUE FACTOR PATHWAY INHIBITOR 2.  
 CC FT DOMAIN 36 86 BPTI/KUNITZ INHIBITOR 1.  
 CC FT DOMAIN 96 149 BPTI/KUNITZ INHIBITOR 2.  
 CC FT DOMAIN 158 208 BPTI/KUNITZ INHIBITOR 3.  
 CC FT DOMAIN 213 217 POLY-LYS.  
 CC FT ACT\_SITE 46 47 REACTIVE BOND (BY SIMILARITY).  
 CC FT ACT\_SITE 107 108 REACTIVE BOND (BY SIMILARITY).  
 CC FT ACT\_SITE 168 169 REACTIVE BOND (BY SIMILARITY).  
 CC FT DISULFID 36 86 REACTIVE BOND (BY SIMILARITY).  
 CC FT DISULFID 45 69 BY SIMILARITY.  
 CC FT DISULFID 61 82 BY SIMILARITY.  
 CC FT DISULFID 96 149 BY SIMILARITY.  
 CC FT DISULFID 106 130 BY SIMILARITY.  
 CC FT DISULFID 122 145 BY SIMILARITY.  
 CC FT DISULFID 158 208 BY SIMILARITY.  
 CC FT DISULFID 167 191 BY SIMILARITY.  
 CC FT DISULFID 183 204 BY SIMILARITY.  
 CC FT CARBOHYD 116 116 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 CC FT CARBOHYD 170 170 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 CC FT CONFLICT 23 23 D->A (IN REF. 4).  
 CC FT SEQUENCE 235 AA: 26934 MW: 975ABA5C53F7C65F CRC64;  
 SO

Query Match 22.6%; Score 214.5; DB 1; Length 235;  
 Best Local Similarity 25.9%; Pred. No. 4.4e-13;  
 Matches 53; Conservative 31; Mismatches 66; Indels 55; Gaps 5;

9 DCLVSKVYVGRASMPRMWYNTDSCGLFYGGCGDGSNNNYLTKKCKK----- 61  
 34 EICLPLDYGPORALILYRYDRTYQSCROFLYGGCGENANNFTWECDDACWRIEYV 93  
 62 -----ATYENATGDL-----ATSRNADSVSPS-----APR 50  
 94 KVCRLQVSDVDCGSESTKTEFFNLSMTCERFFSGGCHNRLENFDPDEATMGFCAPYK 153

QY 90 QSDSDHSSDMFNYEYCTANAVTGPCRASFPWMYDVERNSCNFTYGGCRGNKNSYSE 149  
 DB 154 IPS-----FCYSPDEBLCASNVTRYRPNRYPCDATTYGGGNDNNVSR 201  
 QY 150 EACMLRCF-----RQGNPPPLPLGSK 170  
 DB 202 EDCRRACAKAKRKKMKPKLRFASR 226

RESULT 15  
 APP2\_RAT STANDARD: PRT: 765 AA.  
 AC P15943;  
 DT 01-APR-1990 (Rel. 14, Created)  
 DT 01-OCT-1996 (Rel. 34, last sequence update)  
 DT 01-OCT-2000 (Rel. 40, last annotation update)  
 DE AMYLOID-LIKE PROTEIN 2 PRECURSOR (SPERM MEMBRANE PROTEIN YMK-II).  
 Rattus norvegicus (Rat).  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
 [1]  
 SEQUENCE OF 1-627 FROM N.A.  
 RC STRAIN=WISTAR; TISSUE=BRAIN; AND HEART;  
 RX MEDLINE=94368649; PubMed=8086458;  
 RA Sandbriuk R., Masters C.L., Beyreuther K.;  
 RT "Complete nucleoside and deduced amino acid sequence of rat amyloid  
 RT protein precursor-like protein 2 (APLP2/APPH): two amino acids length  
 RT difference to human and murine homologues.";  
 RL Biochim. Biophys. Acta 1219:167-170(1994).  
 RN [2]  
 RP SEQUENCE OF 575-765 FROM N.A.  
 RC TISSUE=TESTIS;  
 RX MEDLINE=90207205; PubMed=1690887;  
 RA Yan Y.C., Bai Y., Wang L.F., Miao S.Y., Koide S.S.;  
 RT "Characterization of cDNA encoding a human sperm membrane protein  
 RT related to A4 amyloid protein.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 87:2405-2408(1990).  
 CC -1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.  
 CC -1- ALTERNATIVE PRODUCTS: 4 ISOFORMS: A (SHOWN HERE), B, C AND D;  
 CC ARE PRODUCED BY ALTERNATIVE SPLICING.  
 CC -1- SIMILARITY: CONTAINS A PROTEASE INHIBITOR DOMAIN BELONGING TO  
 CC THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 CC -1- SIMILARITY: BELONGS TO THE APP FAMILY.  
 CC -----  
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 CC -----  
 DR EMBL: X77934; CA54906.1;  
 DR EMBL: M31322; AAA42352.1;  
 DR PIR: A35981; A35981.  
 DR HSSP: P05067; ICA0.  
 DR INTERPRO: IPR001868;  
 DR INTERPRO: IPR002223;  
 DR PFWAM: PF00014; Kunitz\_BPTI; 1.  
 DR PRINTS: PR00203; AMYLOIDA4.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00319; A4-EXTRA; 1.  
 DR PROSITE: PS00320; A4-INTRA; 1.  
 DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 1.  
 DR PROSITE: PS00279; BPTI\_KUNITZ\_2; 1.  
 KW Transmembrane; Alternative splicing; Serine protease inhibitor;  
 KW Signal; Glycoprotein.  
 FT SIGNAL 1 29 POTENTIAL.  
 FT CHAIN 30 765 AMYLOID-LIKE PROTEIN 2.  
 FT DOMAIN 30 695 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 696 718 POTENTIAL.

FT DOMAIN 719 765 CYTOPLASMIC (POTENTIAL).  
 FT DOMAIN 218 282 ASP/GLU-RICH (HIGHLY ACIDIC).  
 FT DOMAIN 308 366 BPTI/KUNITZ INHIBITOR.  
 FT ACT\_SITE 322 323 REACTIVE BOND (BY SIMILARITY).  
 FT DISULFID 312 362 BY SIMILARITY.  
 FT DISULFID 321 345 BY SIMILARITY.  
 FT DISULFID 337 358 BY SIMILARITY.  
 FT DOMAIN 218 229 POLY-GLU.  
 FT CARBOHYD 628 628 O-LINKED (GLYCOSAMINOGLYCAN) (POTENTIAL).  
 FT VARSPLIC 311 365 MISSING (IN ISOFORM C AND ISOFORM D).  
 FT VARSPLIC 616 627 MISSING (IN ISOFORM B AND ISOFORM D).  
 FT CONFLICT 575 577 DOF -> ETV (IN REF. 2).  
 SQ SEQUENCE 765 AA; 86882 MM; CF5IFCCCE305A0CF CRC64;

Query Match 22.2%; Score 210; DB 1; Length 765;  
 Best Local Similarity 27.5%; Pred. No. 4,3e-12;  
 Matches 58; Conservative 22; Mismatches 55; Indels 76; Gaps 2;

QY 28 WYVVTGSC-----QLFVYG-----GCDG-----NSNNYLRKE----- 56  
 DB 166 WHIVKAEACLTGNTLYSGMLPCGYDQFHGTEYVCCPQTKVVDSDSTMSKEEEFEE 225  
 QY 57 -----CLKKCATYVENATGDLATSRNAADSVSAPRRQDSE-----DHSSDMFY 102  
 DB 226 DEEDYALDKSEPTT-----ADLEDFTAAADEDEDEEEEGEEVEEDRDYYDSFG 282  
 QY 103 EBY-----CTANAVTGPCRASFPWMYDVERNSCNFTY 130  
 DB 283 DYNENPEPSSDGTISDKREIAHDYKAVSQEAMTGCRVPMRWYFDLSKGRVRFY 302  
 QY 137 GCGRGKNSYRSFEACMLRCFROENPPL 167  
 DB 343 GCGGNNNFESEDDYCAVC--KTMIPPTPL 371

Search completed: January 31, 2001, 15:05:07  
 Job time: 119 sec



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RESULT 5
Q9W004 PRELIMINARY; PRT; 195 AA.
AC Q9W004:
DT 01-NOV-1999 (TREMBLrel. 12, Created)
DT 01-NOV-1999 (TREMBLrel. 12, Last sequence update)
DE HEPATOCYTE GROWTH FACTOR ACTIVATOR INHIBITOR TYPE 2 SPLICE VARIANT 1.
GN HA12.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxId=10090;
RN [1]
RP SEQUENCE FROM N.A.
STRAIN=BA1B/C;
MEDLINE=99160423; PubMed=10049781;
Itoh H., Kataoka H., Hamasuna R., Kitamura N., Kono M.;
RT "Hepatocyte growth factor activator inhibitor type 2 lacking the first
RT Kunitz-type serine proteinase inhibitor domain is a predominant
RT product in mouse but not in human.";
RL Biochem. Biophys. Res. Commun. 255:740-748(1999).
DR EMBL: AF099019; AAD22173.1; -.
DR HSSP: P05067; 1TAW.
DR INTERPRO: IPR002223; -.
DR PFAM: PF00014; Kunitz_BPTI; 1.
DR PRINTS: PR00759; BASICPTASE.
DR PROSITE: PS00280; BPTI_KUNITZ; 1.
KW Serine protease inhibitor.
SQ SEQUENCE 195 AA; 21736 MW; EF49C83AB4E3EDE0 CRC64;

Query Match
Best Local Similarity 40.2%; Score 381; DB 11; Length 195;
Pred. No. 1,7e-33;
Matches 69; Conservative 14; Mismatches 24; Indels 0; Gaps 0;

QY 64 VVENATGDLATSRNADSSVPSAPRRDSDHSDMFNEYECYANAVTGPCRAFPFRMY 123
DB 34 VHEHTDDMARNNMGADSVLSPRKOSADLSAEIFNEYECYPAKAVTGPCRAFPFRMY 93
QY 124 FVERNSCNFNFIYGGCGRGNKNSYRSEACMLRCRPOENPLPLGSK 170
DB 94 YDEKNSCSFTYGGCGRGNKNSLSDCAQCHSGKOMHPLPLPLGK 140

RESULT 6
Q43278 PRELIMINARY; PRT; 513 AA.
AC Q43278:
DT 01-JUN-1998 (TREMBLrel. 06, Created)
DT 01-JUN-1998 (TREMBLrel. 06, Last sequence update)
DE HEPATOCYTE GROWTH FACTOR ACTIVATOR INHIBITOR.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A.
MEDLINE=97197808; PubMed=9045658;
RA Kitamura T., Denda K., Kitamura A., Kawaguchi T., Kito M., Kondo J.,
RA Shimomura T., Takata H., Miyazawa K., Kitamura N.;
RT "Hepatocyte growth factor activator inhibitor, a novel Kunitz-type
RT serine protease inhibitor.";
RL J. Biol. Chem. 272:6370-6376(1997).
DR EMBL: AB000095; BAA25014.1; -.
DR HSSP: P31713; 1SHP.
DR INTERPRO: IPR002172; -.
DR INTERPRO: IPR002223; -.
DR PFAM: PF00014; Kunitz_BPTI; 2.
DR PFAM: PF00057; Idl_recept_a; 1.
DR PFAM: PF00057; Idl_recept_a; 1.
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DR PRINTS: PR00759; BASICPTASE.
DR PROSITE: PS00280; BPTI_KUNITZ; 2.
DR PROSITE: PS01209; IDLRA_1; 1.
DR PROSITE: PS50068; IDLRA_2; 1.
KW Glycoprotein; Serine protease inhibitor.
SQ SEQUENCE 513 AA; 56885 MW; D6E05F3A5885CDDC CRC64;

Query Match
Best Local Similarity 32.9%; Score 312; DB 4; Length 513;
Pred. No. 1,4e-25;
Matches 67; Conservative 24; Mismatches 71; Indels 38; Gaps 3;

QY 5 RSHDFCLVSKYVGRCSRAMPKRWYNTVDGSCQLFYGGCGDSNNYLTRECKLKCATV 7
DB 244 KQEDYCLASNNVGRGSRFPFRMYDPTQICKSFYGGCLGKNNYLTRECKLKCATV 273
QY 65 -----TENATGDLATSRNADSSVPSAPRRDSDHSDMFNEYECYANAVTGPCRAFPFRMY 102
DB 304 QGPMERRHPVCSGTCQPFQFRCNSGCCIDSFLECDTDFPCPDASDEACEKYSQFDEL 363
QY 103 EE-----YCTANAVTGPCRAFPFRMYEDVERNSCNFNFIYGGCGRGNKNSYSEACML 154
DB 364 QIHPSPDKGHCVDLPDTGLCKESIPRWYNPSEHCARFTYGGCYGNKNNFEEDQCLE 423
QY 155 RC-----FROQENPLPL 166
DB 424 SCRGISKVDVFGLRREIRPIP 443

RESULT 7
Q9R097 PRELIMINARY; PRT; 507 AA.
AC Q9R097:
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DE 01-OCT-2000 (TREMBLrel. 15, Last annotation update)
DE HEPATOCYTE GROWTH FACTOR ACTIVATOR INHIBITOR TYPE 1.
GN SPINT1 OR HA11.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxId=10090;
RN [1]
RP SEQUENCE FROM N.A.
STRAIN=BA1B/C;
RA Itoh H., Kataoka H., Kono M.;
RT "Mouse hepatocyte growth factor activator inhibitor type 1 (HA1-1).";
RL Submitted (MAR-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF099018; AAF02490.1; -.
DR HSSP: P05067; 1TAW.
DR MGD: MGI:1338033; Spint1.
DR INTERPRO: IPR002172; -.
DR INTERPRO: IPR002223; -.
DR PFAM: PF00014; Kunitz_BPTI; 2.
DR PFAM: PF00057; Idl_recept_a; 1.
DR PRINTS: PR00759; BASICPTASE.
DR PROSITE: PS00280; BPTI_KUNITZ; 2.
DR PROSITE: PS50068; IDLRA_2; 1.
SQ SEQUENCE 507 AA; 56676 MW; 20CB5DEDCFE46AA7 CRC64;

Query Match
Best Local Similarity 31.0%; Score 294; DB 11; Length 507;
Pred. No. 1,2e-23;
Matches 59; Conservative 23; Mismatches 66; Indels 30; Gaps 3;

QY 9 DFCVSKYVGRCSRAMPKRWYNTVDGSCQLFYGGCGDSNNYLTRECKLKCATV---- 64
DB 242 DYCLASYKVGRCGRSFPFRMYDPTQICKSFYGGCLGKNNYLTRECKLKCATV 301
QY 65 -----TENATGDLATSRNADSSVPSAPRRDSDHSDMFNEYECYANAVTGPCRAFPFRMY 102
DB 302 PKRHVYCSGCHATQRCRSGSIDGFLCEDDTPDPCDGSDEATCEKFTYSGFDELONTH 361
```

	Query Match	27.4%	Score 259.5:	DB 11:	Length 306:
	Best Local Similarity	33.3%	Pred. No. 3.6e-20;		
	Matches	55:	Conservative	21:	Mismatches 64; Indels 25; Gaps 3.
QY	9	DCLVLSKVVGRCASMPKRWNYNTBDSQQLFYVGGDGSNNYLTKEECLKCAT-----	63		
		: : :     : : :     : : :     : : :     : : :     : : :     : : :			
Db	119	DFCFLEEDPGLCGMYKKRFLYNNQTRKOCERFVYGGLGNMNNNETLDECKKICEENVHSP	178		
QY	64	-----YRE-NATGDIATSRNAADSVSAPROPSDDSHSDMEFYEECYANAV	111		
		: : :   : : :   : : :   : : :   : : :   : : :   : : :   : : :			
Db	179	SPVNEYQMSDYVDIGNVTIDRSIVNNIIVVQSFKVERRD-----YGRPRWCLOPAD	230		

Query Match 27.3%; Score 259; DB 5; Length 2230;  
Best Local Similarity 32.9%; Pred. No.3-4e-19;  
Matches 52; Conservative 16; Mismatches 66; Indels 24; Gaps 3;

QY 9 DFCIVSKVYGRKASMPRMWYNTVDGSCQLFYVYGGDGSNNYLKRECKKCAIVTEA 68  
DB 1807 DICEIPAEVGEANYTSMYTDYDQACRFQYVGGCGENEFPEESLACDRKPEPT 1866  
QY 69 TGLAISRMAADSSVSAARRDSDSHSDMNYEECYCANAVTGPCRASFPRWFYDVER 128  
DB 1867 TTTTPARR-----PQPSROD-----VCEDEPAPGDCSTWYLVKWHDRKI 1904  
QY 129 NSCNNEFYGGCRGNKNSYSSEACMLRCRQOENPPLP 166  
DB 1905 GACRCFFYNGCGNGNRFFETENDCCQRCLSQE--PPAP 1940

RESULT 10  
O44938  
ID O44938 PRELIMINARY; PRT; 972 AA.  
AC O44938;  
DT 01-JUN-1998 (TReMBLrel. 06, Created)  
DT 01-JUN-1998 (TReMBLrel. 06, Last sequence update)  
DT 01-OCT-2000 (TReMBLrel. 15, Last annotation update)

DE THROMBOSPONDIN-LIKE PROTEIN (FRAGMENT).  
 GN THRI.  
 OS Haemochus contortus.  
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylida;  
 OC Trichostrongylidae; Haemonchidae; Haemonchinae; Haemonchus.  
 OX NCBI\_TaxID=6289;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=MOREDUN;  
 RA Newlands G.F.J., Skuce P.J.;  
 RL Submitted (JAN-1998) to the EMBL/GenBank/DBJ databases.  
 DR EMBL: AF043121; AAB99830.1; .  
 DR HSSP: P05067; ITAM  
 DR INTERPRO: IPR002223; .  
 DR PFAM: PF00014; Kunitz\_BPTI; 6.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ; 6.  
 DR Serine protease inhibitor.  
 NON\_TER 1  
 SEQUENCE 972 AA; 105301 MW; 4465C6BE8DC19ED CRC64;

Query Match 27.1%; Score 256.5; DB 5; Length 972;  
 Best Local Similarity 30.1%; Pred. No. 2,6e-19;  
 Matches 47; Conservative 21; Mismatches 61; Indels 27; Gaps 1;  
 OY 11 CLVSKVYGRASMPRMWYNTDGSQCLFYVGGCDGNSNNYLTKECLKCATVTENATG 70  
 DB 794 CHLPDVGCGQSFSDSYEMATGSCVEFKYSGSGNANRFSRECENTCV----- 845  
 OY 71 DIATSRNADSSVPSAPRRDSEDSDFNFYEYCTANAVTGPCRASFPFRWYDVER 130  
 DB 846 -----RSEPHSDTSHSTSVCKDEKKEFGCTNATKMYNKKADGT 886  
 OY 131 CNEFTYGGCGKNGKNSYRSEECMLRCFROENPPLP 166  
 DB 887 CNRFHYGCGEGRNFRNEQSCAKANHODACTLP 922

RESULT 11  
 ID 090868 PRELIMINARY; PRT; 3198 AA.  
 AC 090868;  
 DT 01-MAY-2000 (TREMBlrel. 13, Created)  
 DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)  
 DT 01-OCT-2000 (TREMBlrel. 15, Last annotation update)  
 DE LACUNIN PRECURSOR.  
 OS Manduca sexta (Tobacco hawkmoth) (Tobacco hornworm).  
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;  
 OC Phryganea; Neoptera; Endopterygota; Lepidoptera; Glossata; Ditrysia;  
 OC Sphingidae; Sphingidae; Sphinginae; Manduca.  
 OX NCBI\_TaxID=7130;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RA MEDLINE-99457716; PubMed-10528409;  
 RA Nard J.B., Matos R., Walden K.K., Lampe D.J., Robertson H.M.;  
 RT "Expression of lacunin, a large multidomain extracellular matrix  
 protein, accompanies morphogenesis of epithelial monolayers in Manduca  
 sexta".  
 RL Insect Biochem. Mol. Biol. 29:883-897(1999).  
 DR EMBL: AF078161; AAF04457.1; .  
 DR HSSP: P12111; 2KNT.  
 DR INTERPRO: IPR000884; .  
 DR INTERPRO: IPR002221; .  
 DR INTERPRO: IPR002223; .  
 DR INTERPRO: IPR003006; .  
 DR PFAM: PF00014; Kunitz\_BPTI; 10.  
 DR PFAM: PF00047; Ig; 2.  
 DR PFAM: PF00095; wap; 1.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ; 8.  
 DR PROSITE: PS00317; 4\_DISULFIDE\_CORE; 1.  
 KW Signal.

FT SIGNAL 1 21 POTENTIAL.  
 SQ SEQUENCE 3198 AA; 349366 MW; AB4ACD459C0D9134 CRC64;  
 Query Match 26.9%; Score 255; DB 5; Length 3198;  
 Best Local Similarity 31.7%; Pred. No. 1,4e-18;  
 Matches 51; Conservative 23; Mismatches 63; Indels 24; Gaps 3;  
 OY 9 DFLVSKVYGRASMPRMWYNTDGSQCLFYVGGCDGNSNNYLTKECLKCATVTENA 68  
 DB 2133 DLCTLPALIGDCAADYERWYTDIREKSCQRFYGGCAGNNGNFPATQAECCGRC----- 2185  
 OY 69 TGDPLATSRNADSSVPSAPRRDSEDSDFNFYEYCTANAVTGPCRASFPFRWYDVER 128  
 DB 2186 -----SEAKITTVR--PTEAHP-----LTEMCFEMKDPGCTDTRWYDYKL 2228  
 OY 129 NSCNFTYGGCGKNGKNSYRSEECMLRCFROENPPLPUGS 169  
 DB 2229 GKCVTFEYGGCGGNNRNPTEYCYQYCGTADICQLPMRS 2269

RESULT 12  
 ID 093424 PRELIMINARY; PRT; 287 AA.  
 AC 093424;  
 DT 01-NOV-1998 (TREMBlrel. 08, Created)  
 DT 01-NOV-1998 (TREMBlrel. 08, Last sequence update)  
 DT 01-JUN-2000 (TREMBlrel. 14, Last annotation update)  
 DE HYPOTHETICAL 33.1 KDA PROTEIN.  
 OS Cyprinus carpio (Common carp).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Ostariophysi;  
 OC Cypriniformes; Cyprinidae; Cyprininae; Cyprinus.  
 OX NCBI\_TaxID=7962;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=LIVER;  
 RA Gracey A.Y.;  
 RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.  
 DR EMBL: AF008648; AAC19410.1; .  
 DR HSSP: P31713; 1SHP.  
 DR INTERPRO: IPR002223; .  
 DR PFAM: PF00014; Kunitz\_BPTI; 3.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ; 3.  
 KW Hypothetical protein; Serine protease inhibitor.  
 SQ SEQUENCE 287 AA; 33093 MW; DF69B3D76718115E CRC64;

Query Match 26.6%; Score 252; DB 13; Length 287;  
 Best Local Similarity 30.9%; Pred. No. 2,1e-19;  
 Matches 50; Conservative 19; Mismatches 53; Indels 40; Gaps 3;  
 OY 8 HDCLVSKVYGRASMPRMWYNTDGSQCLFYVGGCDGNSNNYLTKECLKCATVTEN 67  
 DB 39 HHSCLAKDKGPKAKLDREYFDIDGRCESEFYGGCGQGNENFETLQCEKKCLV---- 94  
 OY 68 ATGDPLATSRNADSSVPSAPRRDSEDSDFNFYEYCTANAVTGPCRASFPFRWYDVER 127  
 DB 95 -----KDKKSP-----CQLDDEPQPCGVLPRFYDFDK 122  
 OY 128 NSCNFTYGGCGKNGKNSYRSEECMLRCF---ROENPPL 165  
 DB 123 SQCKRFFYGGCGGNANRNPTEYCYQYCGTADICQLPMRS 2269

RESULT 13  
 ID 076840 PRELIMINARY; PRT; 2167 AA.  
 AC 076840; Q22911;  
 DT 01-NOV-1998 (TREMBlrel. 08, Created)  
 DT 01-NOV-1998 (TREMBlrel. 08, Last sequence update)  
 DT 01-JUN-2000 (TREMBlrel. 14, Last annotation update)









QY 121 RMYEDVERNSCNNTFYGGCGRNKNSYSEACMLRCFROQENPPLPGSK 170  
 DB 148 RMYEDVERNSCNNTFYGGCGRNKNSYSEACMLRCFROQENPPLPGSK 197

## RESULT 2

000271 PRELIMINARY: PRT: 252 AA.

AC 000271: PRELIMINARY: PRT: 252 AA.  
 DT 01-JUL-1997 (TREMBLrel. 04, Created)  
 DT 01-JUL-1997 (TREMBLrel. 04, Last sequence update)  
 DT 01-OCT-2000 (TREMBLrel. 15, Last annotation update)  
 DE BIKUNIN.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
 OX NCBI\_TaxID=9606;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RA TISSUE=PLACENTA;  
 RA MEDLINE=9727372; PubMed=9115294;  
 RA Marlor C.W., Delaria K.A., Davis G., Muller D.K., Greve J.M.,  
 RA Tamburini P.P.;  
 RA "Identification and cloning of human placental bikunin, a novel serine  
 RA protease inhibitor containing two Kunitz domains.";  
 RA J. Biol. Chem. 272:12202-12208(1997).  
 DR HSSP: P05067; ITAM  
 DR HSSP: P05067; ITAM  
 DR INTERPRO: IPR002223;  
 DR PFAM: PF00014; Kunitz\_BPTI; 2.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ; 2.  
 KW Serine protease inhibitor.  
 SQ SEQUENCE 252 AA; 28228 MW; A7D3360C0EBCAB2B CRC64;

Query Match 100.0%; Score 948; DB 4; Length 252;  
 Best Local Similarity 100.0%; Pred. No. 1.7e-94;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADDRESSHDFCLVSKYVGRCRASMPRMWYNTDSCQLFYGGCGDGNNSNYLTKKECLTK 60  
 DB 28 ADDRESSHDFCLVSKYVGRCRASMPRMWYNTDSCQLFYGGCGDGNNSNYLTKKECLTK 87  
 QY 61 CATVTENATGDLATSRNADSSVPSAPRRDSEHSSDMFNYEYCTANAVTGCRASFP 120  
 DB 88 CATVTENATGDLATSRNADSSVPSAPRRDSEHSSDMFNYEYCTANAVTGCRASFP 147  
 DB 121 RMYEDVERNSCNNTFYGGCGRNKNSYSEACMLRCFROQENPPLPGSK 170  
 DB 148 RMYEDVERNSCNNTFYGGCGRNKNSYSEACMLRCFROQENPPLPGSK 197

DR HSSP: P05067; ITAM.  
 DR INTERPRO: IPR002223;  
 DR PFAM: PF00014; Kunitz\_BPTI; 2.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ; 2.  
 KW Protease; Serine protease inhibitor.  
 SQ SEQUENCE 252 AA; 28231 MW; B21593466413841E CRC64;

Query Match 99.7%; Score 945; DB 4; Length 252;  
 Best Local Similarity 99.4%; Pred. No. 3.3e-94;  
 Matches 169; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADDRESSHDFCLVSKYVGRCRASMPRMWYNTDSCQLFYGGCGDGNNSNYLTKKECLTK 60  
 DB 28 ADDRESSHDFCLVSKYVGRCRASMPRMWYNTDSCQLFYGGCGDGNNSNYLTKKECLTK 87  
 QY 61 CATVTENATGDLATSRNADSSVPSAPRRDSEHSSDMFNYEYCTANAVTGCRASFP 120  
 DB 88 CATVTENATGDLATSRNADSSVPSAPRRDSEHSSDMFNYEYCTANAVTGCRASFP 147  
 QY 121 RMYEDVERNSCNNTFYGGCGRNKNSYSEACMLRCFROQENPPLPGSK 170  
 DB 148 RMYEDVERNSCNNTFYGGCGRNKNSYSEACMLRCFROQENPPLPGSK 197

## RESULT 4

09WU03 PRELIMINARY: PRT: 252 AA.

AC 09WU03: PRELIMINARY: PRT: 252 AA.  
 DT 01-NOV-1999 (TREMBLrel. 12, Created)  
 DT 01-NOV-1999 (TREMBLrel. 12, Last sequence update)  
 DT 01-OCT-2000 (TREMBLrel. 15, Last annotation update)  
 DE HEPATOCYTE GROWTH FACTOR ACTIVATOR INHIBITOR TYPE 2.  
 GN HA12.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 OX NCBI\_TaxID=10096;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RA STRAIN=BA1B/C;  
 RA MEDLINE=99160423; PubMed=10049781;  
 RA Itoh H., Kataoka H., Hamasuna R., Kitamura N., Koono M.;  
 RA Hepatocyte growth factor activator inhibitor type 2 lacking the first  
 RA Kunitz-type serine protease inhibitor domain is a predominant  
 RT product in mouse but not in human.";  
 RL Biochem. Biophys. Res. Commun. 255:740-748(1999).  
 DR HSSP: P05067; ITAM.  
 DR HSSP: P05067; ITAM.  
 DR INTERPRO: IPR002223;  
 DR PFAM: PF00014; Kunitz\_BPTI; 2.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ; 2.  
 KW Serine protease inhibitor.  
 SQ SEQUENCE 252 AA; 27914 MW; B2FF4B86924D4F8F CRC64;

Query Match 69.9%; Score 663; DB 11; Length 252;  
 Best Local Similarity 68.2%; Pred. No. 9e-64;  
 Matches 116; Conservative 21; Mismatches 33; Indels 0; Gaps 0;

QY 1 ADDRESSHDFCLVSKYVGRCRASMPRMWYNTDSCQLFYGGCGDGNNSNYLTKKECLTK 60  
 DB 28 ASRELVDHESGVSKYVGRCRASIPRMWYNTDSCQPFYGGCGDGNNSYOSKECLTK 87  
 QY 61 CATVTENATGDLATSRNADSSVPSAPRRDSEHSSDMFNYEYCTANAVTGCRASFP 120  
 DB 88 CAGVTENTTDMANRNADSSVPSAPRRDSEHSSDMFNYEYCTANAVTGCRASFP 147  
 QY 121 RMYEDVERNSCNNTFYGGCGRNKNSYSEACMLRCFROQENPPLPGSK 170  
 DB 148 RMYEDVERNSCNNTFYGGCGRNKNSYSEACMLRCFROQENPPLPGSK 197

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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:02:44 ; Search time 23.18 seconds  
(without alignments)  
859.592 Million cell updates/sec

Title: US-09-441-654A-1

Perfect score: 948  
Sequence: 1 ADREKSHDFCLVSKVYGRG.....ACMLRCFRQENPPLIGSK 170

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 374700 seqs, 117207915 residues

total number of hits satisfying chosen parameters: 374700

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database :  
1: SP archaea:\*  
2: SP bacteria:\*  
3: SP fungi:\*  
4: SP human:\*  
5: SP invertebrate:\*  
6: SP mammal:\*  
7: SP mnc:\*  
8: SP organelle:\*  
9: SP phage:\*  
10: SP plant:\*  
11: SP rodent:\*  
12: SP virus:\*  
13: SP vertebrate:\*  
14: SP unclassified:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	948	100.0	252	4	043291 homo sapien
2	948	100.0	252	4	000271 homo sapien
3	945	99.7	252	4	014895 homo sapien
4	663	69.9	252	11	09WU03 mus musculu
5	381	40.2	195	11	09WU04 mus musculu
6	312	32.9	513	4	043278 homo sapien
7	294	31.0	507	11	09R097 mus musculu
8	259.5	27.4	306	11	054819 mus musculu
9	259	27.3	2230	5	09VAV4 drosophila
10	256.5	27.1	972	5	044938 haemochnus
11	255	26.9	3198	5	09U868 manduca sex
12	252	26.6	287	13	093424 cyprinus ca
13	250.5	26.4	2167	5	076840 caenorhabdi
14	244.5	25.8	151	4	078491 homo sapien
15	240	25.3	2225	5	045881 caenorhabdi
16	239.5	25.3	396	6	028874 canis famli
17	235.5	24.8	144	11	090W87 mesocricetu
18	233.5	24.6	352	11	070160 cavia porce
19	228	24.1	251	4	095103 homo sapien

20	227.5	24.0	246	11	092208 mus musculu
21	222	23.4	1043	5	017644 caenorhabdi
22	219.5	23.2	342	13	P70004 xenopus lae
23	219	23.1	922	5	021418 caenorhabdi
24	209	22.0	1743	5	09XW85 caenorhabdi
25	208	21.9	751	11	060709 mus musculu
26	208	21.9	763	11	061482 mus musculu
27	204.5	21.6	230	11	035536 mus musculu
28	200	21.1	523	4	014594 mus musculu
29	200	21.1	1599	5	009983 mus musculu
30	199	21.0	1522	5	022685 caenorhabdi
31	198	20.9	1195	5	09N343 caenorhabdi
32	195	20.6	1391	5	019021 caenorhabdi
33	194	20.5	1297	5	09U350 caenorhabdi
34	193	20.4	1474	5	062504 caenorhabdi
35	186.5	19.7	747	13	091963 xenopus lae
36	186	19.6	484	4	013793 homo sapien
37	186	19.6	547	4	013764 homo sapien
38	186	19.6	770	6	09T010 sus scrofa
39	185.5	19.6	1203	5	045916 caenorhabdi
40	183	19.3	160	11	09Q278 caenorhabdi
41	178	18.8	1965	5	061893 caenorhabdi
42	177	18.7	59	5	09TWF8 anemonia s-
43	174.5	18.4	692	5	045101 caenorhabdi
44	174.5	18.4	780	13	073683 tetracoda
45	173	18.2	58	5	09TWF9 anemonia eu

## ALIGNMENTS

RESULT	ID	SEQUENCE FROM N.A.	PRELIMINARY:	PRT:	252 AA.
043291	043291	01-JUN-1998 (TRENBLREL. 06, Created)			
AC	043291	01-JUN-1998 (TRENBLREL. 06, Last sequence update)			
DT	01-JUN-1998	01-OCT-2000 (TRENBLREL. 15, Last annotation update)			
DT	01-OCT-2000	HEPATOCYTE GROWTH FACTOR ACTIVATOR INHIBITOR TYPE 2.			
DE	HEPATOCYTE GROWTH FACTOR ACTIVATOR INHIBITOR TYPE 2.				
OC	Homo sapiens (Human).				
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;				
OC	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
OX	NCBI_TaxID=9606;				
RN	[1]	SEQUENCE FROM N.A.			
RP	MEDLINE=96010584; PubMed=9346890;				
RX	Kawaguchi T., Qin L., Shimomura T., Kondo J., Matsumoto K., Denta K.,				
RA	Kitamura N.;				
RT	"Purification and cloning of hepatocyte growth factor activator				
RT	inhibitor type 2, a Kunitz-type serine protease inhibitor.";				
RL	J. Biol. Chem. 272:27558-27564(1997).				
DR	EMBL; AB006534; BAA25024.1; -				
DR	HSSP; P05067; ITAM.				
DR	INTERPRO: IPR002223; -				
DR	PFAM: PF00014; Kunitz_BPTI; 2.				
DR	PRINTS: PR00759; BASICPTASE.				
DR	PROSITE: PS00280; BPTI_KUNITZ; 2.				
KW	Serine protease inhibitor.				
SQ	SEQUENCE 252 AA; 28169 MW; F7D3D834ED631DF0 CRC64;				

Query Match 100.0%; Score 948; DB 4; Length 252;  
Best Local Similarity 100.0%; Pred. No. 1.7e-94;  
Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	1	ADREKSHDFCLVSKVYGRGASPRWYVNTGSCOLFYYGGGDSNNYLFKECLCK 60
DB	28	ADREKSHDFCLVSKVYGRGASPRWYVNTGSCOLFYYGGGDSNNYLFKECLCK 87
QY	61	CATYENATGDLATSRNADSVSPAPRRDSEHSSDMRYEYCTANAVTGCRASFP 120
DB	88	CATYENATGDLATSRNADSVSPAPRRDSEHSSDMRYEYCTANAVTGCRASFP 147

GenCore version 4.5  
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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:02:39 ; Search time 16.59 seconds

(without alignments)  
350,388 Million cell updates/sec

Title: US-09-441-654a-1

Sequence: 948  
I ADRRSIHDECLVSKVVGRC.....ACMDRCFROENPLPLGSK 170

Scoring table:

BLOSUM62  
Gap 10.0 , Gapext 0.5

Searched: 268485 seqs, 34193795 residues

Minimum number of hits satisfying chosen parameters: 268485

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database :

A\_Geneseq\_36:\*

- 1: /SIDSL/gcgdata/geneseq/geneseqp/AA1980.DAT:\*
- 2: /SIDSL/gcgdata/geneseq/geneseqp/AA1981.DAT:\*
- 3: /SIDSL/gcgdata/geneseq/geneseqp/AA1982.DAT:\*
- 4: /SIDSL/gcgdata/geneseq/geneseqp/AA1983.DAT:\*
- 5: /SIDSL/gcgdata/geneseq/geneseqp/AA1984.DAT:\*
- 6: /SIDSL/gcgdata/geneseq/geneseqp/AA1985.DAT:\*
- 7: /SIDSL/gcgdata/geneseq/geneseqp/AA1986.DAT:\*
- 8: /SIDSL/gcgdata/geneseq/geneseqp/AA1987.DAT:\*
- 9: /SIDSL/gcgdata/geneseq/geneseqp/AA1988.DAT:\*
- 10: /SIDSL/gcgdata/geneseq/geneseqp/AA1989.DAT:\*
- 11: /SIDSL/gcgdata/geneseq/geneseqp/AA1990.DAT:\*
- 12: /SIDSL/gcgdata/geneseq/geneseqp/AA1991.DAT:\*
- 13: /SIDSL/gcgdata/geneseq/geneseqp/AA1992.DAT:\*
- 14: /SIDSL/gcgdata/geneseq/geneseqp/AA1993.DAT:\*
- 15: /SIDSL/gcgdata/geneseq/geneseqp/AA1994.DAT:\*
- 16: /SIDSL/gcgdata/geneseq/geneseqp/AA1995.DAT:\*
- 17: /SIDSL/gcgdata/geneseq/geneseqp/AA1996.DAT:\*
- 18: /SIDSL/gcgdata/geneseq/geneseqp/AA1997.DAT:\*
- 19: /SIDSL/gcgdata/geneseq/geneseqp/AA1998.DAT:\*
- 20: /SIDSL/gcgdata/geneseq/geneseqp/AA1999.DAT:\*
- 21: /SIDSL/gcgdata/geneseq/geneseqp/AA2000.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	948	100.0	170	W30041	Human placental bi
2	948	100.0	179	W30053	Human placental bi
3	948	100.0	197	W30043	Human placental bi
4	948	100.0	213	W30042	Human placental bi
5	948	100.0	225	W30046	Human placental bi
6	948	100.0	235	W30060	Human placental bi
7	948	100.0	240	W30045	Human placental bi
8	948	100.0	248	W30044	Human placental bi
9	948	100.0	252	W30040	Human placental bi
10	948	100.0	252	W30040	Human placental bi
11	948	100.0	252	W13665	Hepatocyte growth
12	859	90.6	153	W30051	Human placental bi

13	819	86.4	146	W30052	Human placental bi
14	750	79.1	170	W30061	Human placental bi
15	501	52.8	92	W30054	Human placental bi
16	488	51.5	130	W30062	EST R35464 protein
17	487	51.4	169	W30063	EST R74593 protein
18	337	35.5	58	W30049	Human placental bi
19	334	35.2	58	W30049	Human placental bi
20	312	32.9	513	W30047	Hepatocyte growth
21	312	32.9	513	W30047	Human placental bi
22	297	31.3	51	W30048	Human placental bi
23	297	31.3	51	W30048	Human placental bi
24	281	29.6	128	W30050	Human placental bi
25	277	29.2	128	W30050	Human placental bi
26	276	29.1	128	W30050	Human placental bi
27	274	28.9	128	W30050	Human placental bi
28	273	28.8	128	W30050	Human placental bi
29	272.5	28.7	124	W30050	Human placental bi
30	272.5	28.7	144	W30050	Human placental bi
31	272	28.7	128	W30050	Human placental bi
32	272	28.7	128	W30050	Human placental bi
33	272	28.7	128	W30050	Human placental bi
34	271.5	28.6	124	W30050	Human placental bi
35	271.5	28.6	144	W30050	Human placental bi
36	271.5	28.6	145	W30050	Human placental bi
37	271.5	28.6	145	W30050	Human placental bi
38	267	28.2	128	W30050	Human placental bi
39	267	28.2	128	W30050	Human placental bi
40	266.5	28.1	128	W30050	Human placental bi
41	265	28.0	128	W30050	Human placental bi
42	261	27.5	128	W30050	Human placental bi
43	259	27.3	128	W30050	Human placental bi
44	255	26.9	128	W30050	Human placental bi
45	254.5	26.8	124	W30050	Human placental bi

#### ALIGNMENTS

RESULT	1	
ID	W30041	Standard; Protein; 170 AA.
XX	W30041;	
XX	W30041;	
XX	20-APR-1998 (first entry)	
XX	Human placental, bikunin.	
XX	Human; placental bikunin; inhibition; trypsin; kallikrein;	
KW	plasma; factor xila; treatment; prevention; oedema;	
KW	inflammation; infection; granulomatosis; multiple sclerosis;	
KW	ischaemia; peroperative blood loss; sepsis; shock; fibrosis;	
KW	blood coagulation disease; polytrauma; stroke; haemorrhage;	
XX	gastric cancer; cervical cancer; metastasis; blood loss.	
XX	Homo sapiens.	
OS	Homo sapiens.	
XX	WO9733996-A2.	
XX	18-SEP-1997.	
XX	10-MAR-1997;	97WO-US03894.
XX	04-OCT-1996;	96US-0725251.
XX	11-MAR-1996;	96US-0013106.
XX	14-JUN-1996;	96US-0019793.
XX	(FARB ) BAYER CORP.	
XX	Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;	
XX	WPI; 1997-470876/43.	

PR New human placental bikunin - used to inhibit kallikrein, trypsin  
PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
PT perioperative blood loss

XX Claim 1: Page 65; 110pp; English.

XX The present sequence is a human placental bikunin, which  
CC inhibits, e.g. trypsin, kallikrein, plasmin and factor X11a.  
CC Bikunin can be used to treat or prevent brain and spinal cord  
CC oedema, inflammation, infection or granulomatosis, multiple  
CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
CC cerebral or subarachnoid haemorrhage and gastric or cervical  
CC cancer and prevent metastasis. It is particularly useful for  
CC reducing blood loss during surgery, and can also be used to treat  
CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
CC influenza and similar viral infections, acute pancreatitis and  
CC gout, and prevent pre-term labour. It has similar properties to  
CC apoferritin, but is less highly charged so should be less  
CC immunogenic and less likely to damage the kidneys. Manipulation  
CC of the bikunin sequence may allow the inhibitory profile to be  
CC altered. It also reduces or eliminates the need for whole donor  
CC blood or blood products during surgery, thereby reducing the risk  
CC of infection and other adverse side effects, as well as reducing  
CC the cost of surgery.

XX Sequence 170 AA;

Query Match 100.0%; Score 948; DB 18; Length 170;  
Best Local Similarity 100.0%; Pred. No. 1.2e-89;  
Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADDRESSHDFCLVSVYVGRCRASMPRMWYNTDGSQCLFVYGGCDGNSNNYLTKKECLK 60  
DB 1 addresshdfclvsvygrcrasmprrwvnyvdgscqlfvygdcgnsnnyltkkeclk 60

QY 61 CATVTENATGDIATSRNAADSSVPSAPRRQDSEHSDMFNEEYECTANAVGPCRASFP 120  
DB 61 catvtenatgdiatsrnaadssvpsaprrqdsedhsdmfneyectanavgpcrafp 120

QY 121 RWYFDVERNSCNFTYGGCRGNKNSYSEACMLRCFROENPPLPLGSK 170  
DB 121 rwyfdvernschnftlyggcrgnknsyrseacmlrcfrgenpplplgsk 170

RESULT 2  
W30053 standard; Protein: 179 AA.

W30053;

20-APR-1998 (first entry)

Human placental bikunin.

XX Human; placental bikunin; inhibition: trypsin; kallikrein;  
XX plasmin; factor X11a; treatment: prevention; oedema;  
XX inflammation; infection; granulomatosis; multiple sclerosis;  
XX ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
XX blood coagulation disease; polytrauma; stroke; haemorrhage;  
XX gastric cancer; cervical cancer; metastasis; blood loss.

XX Homo sapiens.

XX WO9733996-A2.

XX 18-SEP-1997.

XX 10-MAR-1997; 97WO-US03894.

XX 04-OCT-1996; 96US-0725251.  
XX 11-MAR-1996; 96US-0013106.

PR 14-JUN-1996; 96US-0019793.

XX (FARB ) BAYER CORP.

XX Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;

XX WPI; 1997-470876/43.

PR New human placental bikunin - used to inhibit kallikrein, trypsin  
PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
PT perioperative blood loss

XX Claim 1: Page 67; 110pp; English.

XX The present sequence is a human placental bikunin, which  
CC inhibits, e.g. trypsin, kallikrein, plasmin and factor X11a.  
CC Bikunin can be used to treat or prevent brain and spinal cord  
CC oedema, inflammation, infection or granulomatosis, multiple  
CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
CC cerebral or subarachnoid haemorrhage and gastric or cervical  
CC cancer and prevent metastasis. It is particularly useful for  
CC reducing blood loss during surgery, and can also be used to treat  
CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
CC influenza and similar viral infections, acute pancreatitis and  
CC gout, and prevent pre-term labour. It has similar properties to  
CC apoferritin, but is less highly charged so should be less  
CC immunogenic and less likely to damage the kidneys. Manipulation  
CC of the bikunin sequence may allow the inhibitory profile to be  
CC altered. It also reduces or eliminates the need for whole donor  
CC blood or blood products during surgery, thereby reducing the risk  
CC of infection and other adverse side effects, as well as reducing  
CC the cost of surgery.

XX Sequence 179 AA;

Query Match 100.0%; Score 948; DB 18; Length 179;  
Best Local Similarity 100.0%; Pred. No. 1.3e-89;  
Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADDRESSHDFCLVSVYVGRCRASMPRMWYNTDGSQCLFVYGGCDGNSNNYLTKKECLK 60  
DB 1 addresshdfclvsvygrcrasmprrwvnyvdgscqlfvygdcgnsnnyltkkeclk 60

QY 61 CATVTENATGDIATSRNAADSSVPSAPRRQDSEHSDMFNEEYECTANAVGPCRASFP 120  
DB 61 catvtenatgdiatsrnaadssvpsaprrqdsedhsdmfneyectanavgpcrafp 120

QY 121 RWYFDVERNSCNFTYGGCRGNKNSYSEACMLRCFROENPPLPLGSK 170  
DB 121 rwyfdvernschnftlyggcrgnknsyrseacmlrcfrgenpplplgsk 170

RESULT 3  
W30043 standard; Protein: 197 AA.

W30043;

20-APR-1998 (first entry)

Human placental bikunin.

XX Human; placental bikunin; inhibition: trypsin; kallikrein;  
XX plasmin; factor X11a; treatment: prevention; oedema;  
XX inflammation; infection; granulomatosis; multiple sclerosis;  
XX ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
XX blood coagulation disease; polytrauma; stroke; haemorrhage;  
XX gastric cancer; cervical cancer; metastasis; blood loss.

XX Homo sapiens.



PN W09733996-A2.  
 XX 18-SEP-1997.  
 XX 10-MAR-1997; 97WO-US03894.  
 XX 04-OCT-1996; 96US-0725251.  
 PR 11-MAR-1996; 96US-0013106.  
 PR 14-JUN-1996; 96US-0019793.  
 XX (FARB ) BAYER CORP.  
 PA Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 PI WPI; 1997-470876/43.  
 DR New human placental bikunin - used to inhibit kallikrein, trypsin  
 XX etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 XX perioperative blood loss  
 XX Claim 1; Page 65; 110pp; English.  
 CC The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIIa.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.  
 CC Sequence 197 AA;  
 SQ

Query Match 100.0%; Score 948; DB 18; Length 197;  
 Best Local Similarity 100.0%; Pred. No. 1.4e-89;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 1 ADDRESSHDFCLVSKVGRASMPRMWYNTDGSQCLFYGGCDGNSNNYLTKEECLK 60  
 19 addresshdfclvskvgrasmprrwvnyntdgsqclfyggcdgnsnnyltkeecik 78  
 QY 61 CATVTENATGDLATSRNAADSSVPSAPRRDSEHSSDMFYEYCTANAVTGPCRASFP 120  
 DB 79 catvtenatgdlatsrnaadssvpsaprrqdsedhsdmtfyeyctanaavtgcrafp 138  
 QY 121 RWFEDVRNSCNMFYGGCGKNKNSYRSEACMLRCFROENPPLPGSK 170  
 DB 139 rwyfdvnrscnmfiyggcgknksyrseacmlrcfrqenpplpgsk 188

RESULT 4  
 W30042  
 ID W30042 standard; Protein; 213 AA.  
 XX W30042;  
 AC  
 XX 20-APR-1998 (first entry)  
 DT Human placental bikunin.  
 XX Human placental bikunin.  
 DE  
 XX Human; placental bikunin; inhibition; trypsin; kallikrein;  
 KW

KW Plasmin; factor XIIIa; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; hemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.  
 XX Homo sapiens.  
 XX W09733996-A2.  
 XX 18-SEP-1997.  
 XX 10-MAR-1997; 97WO-US03894.  
 XX 04-OCT-1996; 96US-0725251.  
 PR 11-MAR-1996; 96US-0013106.  
 PR 14-JUN-1996; 96US-0019793.  
 XX (FARB ) BAYER CORP.  
 PA Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 PI WPI; 1997-470876/43.  
 DR New human placental bikunin - used to inhibit kallikrein, trypsin  
 XX etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 XX perioperative blood loss  
 XX Claim 1; Page 65; 110pp; English.  
 CC The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIIa.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.  
 CC Sequence 213 AA;  
 SQ

Query Match 100.0%; Score 948; DB 18; Length 213;  
 Best Local Similarity 100.0%; Pred. No. 1.6e-89;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 1 ADDRESSHDFCLVSKVGRASMPRMWYNTDGSQCLFYGGCDGNSNNYLTKEECLK 60  
 1 addresshdfclvskvgrasmprrwvnyntdgsqclfyggcdgnsnnyltkeecik 78  
 QY 61 CATVTENATGDLATSRNAADSSVPSAPRRDSEHSSDMFYEYCTANAVTGPCRASFP 120  
 DB 61 catvtenatgdlatsrnaadssvpsaprrqdsedhsdmtfyeyctanaavtgcrafp 138  
 QY 121 RWFEDVRNSCNMFYGGCGKNKNSYRSEACMLRCFROENPPLPGSK 170  
 DB 121 rwyfdvnrscnmfiyggcgknksyrseacmlrcfrqenpplpgsk 170

RESULT 5  
 W30046  
 ID W30046 standard; Protein; 225 AA.  
 KW

XX W30046;  
 XX 20-APR-1998 (first entry)  
 DT  
 XX  
 DE Human placental bikunin.  
 XX  
 KW Human: Placental bikunin; inhibition; trypsin; kallikrein;  
 KW plasmin; factor XIIa; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.  
 OS  
 XX Homo sapiens.  
 XX  
 PN W09733996-A2.  
 XX  
 PD 18-SEP-1997.  
 XX  
 X 10-MAR-1997; 97WO-US03894.  
 XX  
 PR 04-OCT-1996; 96US-0725251.  
 PR 11-MAR-1996; 96US-0013106.  
 PR 14-JUN-1996; 96US-0019793.  
 PA (FARB ) BAYER CORP.  
 XX  
 PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 XX  
 XX WPI: 1997-470876/43.  
 XX  
 PT New human placental bikunin - used to inhibit kallikrein, trypsin  
 PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 PT perioperative blood loss  
 PS  
 XX  
 XX Claim 1; Page 66; 110pp; English.  
 XX  
 CC The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC apocitin, but is less likely to damage the kidneys. Manipulation  
 CC immunogenic and less likely to damage the inhibitory profile to be  
 CC of the bikunin sequence may allow the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.  
 CC  
 CC Sequence 225 AA;  
 SQ  
 Query Match 100.0%; Score 948; DB 18; Length 225;  
 Best Local Similarity 100.0%; Pred. No. 1.7e-89;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 1 ADDRESSIDFCLVSKVRCRASRSMFRMYNTYDSCQFLYGGCGGNSNNYTKKECLKK 60  
 DB 1 addressidfclyskvrcrasrsmfrmyntydgscqfltyggcggnsmnytkkeclkk 60  
 OY 61 CATVENATGDLATSRNADSSVSPAPRRDSEDSMDENYETECITANAVTGPCRASFP 120  
 DB 61 catvenatgdlatsrnaadssvspaprrdseedsmdenfyeyctanavtgpccrasfp 120

OY 121 RMYEDVERNSCNNTYGGCRGNKNSYRSEACMLRCFROENPPLPLGSK 170  
 DB 121 rmyedvernschnfiyggcrgnknsyrseacmlrcfrqenpplpplgsk 170  
 RESULT 6  
 W30060  
 ID W30060 standard; Protein; 235 AA.  
 XX  
 AC W30060;  
 XX  
 DT 20-APR-1998 (first entry)  
 XX  
 DE Human consensus bikunin.  
 XX  
 KW Human: consensus bikunin; inhibition; trypsin; kallikrein;  
 KW plasmin; factor XIIa; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.  
 OS  
 XX Homo sapiens.  
 XX  
 PN W09733996-A2.  
 XX  
 PD 18-SEP-1997.  
 XX  
 PF 10-MAR-1997; 97WO-US03894.  
 XX  
 PR 04-OCT-1996; 96US-0725251.  
 PR 11-MAR-1996; 96US-0013106.  
 PR 14-JUN-1996; 96US-0019793.  
 PA (FARB ) BAYER CORP.  
 XX  
 PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 XX  
 XX WPI: 1997-470876/43.  
 XX  
 DR N-PSDB; T90732.  
 XX  
 PT New human placental bikunin - used to inhibit kallikrein, trypsin  
 PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 PT perioperative blood loss  
 PS  
 XX Disclosure; Fig 3; 110pp; English.  
 XX  
 CC The present sequence is a consensus human bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation disease, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC apocitin, but is less likely to damage the kidneys. Manipulation  
 CC immunogenic and less likely to damage the inhibitory profile to be  
 CC of the bikunin sequence may allow the need for whole donor  
 CC altered. It also reduces or eliminates the need for whole donor

CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.  
 CC  
 XX  
 SQ Sequence 235 AA;

Query Match 100.0%; Score 948; DB 18; Length 235;  
 Best Local Similarity 100.0%; Pred. No. 1.8e-89;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADRESIHDFCLVSKVYGRASMPRWYNTDSCQLFYVGGCDGNSNNTLKEECLK 60  
 Db 19 adrehsidfcivskvgrcrasmprrwvntdgsqclfyvgcdgnsnnyltkeecik 78  
 QY 61 CATVTENATGDIATSRNAADSVSPAPRRDSEHSSDMFYEEYCTANAVTGFCRASFP 120  
 Db 79 catvtenatgdiatsrnaadsvspaprrdsehdssdmfyeyctanavtgcrafp 138  
 121 RWFEDVERNSCNFFIYGGCGRGNKNSYRSEBACMLRCFROENPPLPGSK 170  
 139 rwyfdivernscnffiyggcgrgnknsyrseacmlrcfrqenpplpgsk 188

RESULT 7  
 W30045  
 ID W30045 standard; Protein; 240 AA.  
 AC W30045;

DT 20-APR-1998 (first entry)  
 DE Human placental bikunin.

Human; placental bikunin; inhibition; trypsin; kallikrein;  
 plasmin; factor XIIa; treatment; prevention; oedema;  
 inflammation; infection; granulomatosis; multiple sclerosis;  
 ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 blood coagulation disease; polytrauma; stroke; haemorrhage;  
 gastric cancer; cervical cancer; metastasis; blood loss.

OS Homo sapiens.  
 XX  
 PN WO9733996-A2.  
 XX  
 PD 18-SEP-1997.  
 XX  
 10-MAR-1997; 97WO-US03894.  
 04-OCT-1996; 96US-0725251.  
 11-MAR-1996; 96US-0013106.  
 14-JUN-1996; 96US-0019793.

(FARB ) BAYER CORP.

PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 DR N-PSDB; T90734.  
 N-PSDB; T90734.

PT New human placental bikunin - used to inhibit kallikrein, trypsin  
 PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 PT perioperative blood loss

PS Claim 1; Page 66; 110pp; English.

CC The present sequence is human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
 CC bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical

CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.  
 CC  
 XX  
 SQ Sequence 240 AA;

Query Match 100.0%; Score 948; DB 18; Length 240;  
 Best Local Similarity 100.0%; Pred. No. 1.8e-89;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADRESIHDFCLVSKVYGRCRASMPRWYNTDSCQLFYVGGCDGNSNNTLKEECLK 60  
 Db 28 adrehsidfcivskvgrcrasmprrwvntdgsqclfyvgcdgnsnnyltkeecik 67  
 QY 61 CATVTENATGDIATSRNAADSVSPAPRRDSEHSSDMFYEEYCTANAVTGFCRASFP 120  
 Db 88 catvtenatgdiatsrnaadsvspaprrdsehdssdmfyeyctanavtgcrafp 147  
 QY 121 RWFEDVERNSCNFFIYGGCGRGNKNSYRSEBACMLRCFROENPPLPGSK 170  
 148 rwyfdivernscnffiyggcgrgnknsyrseacmlrcfrqenpplpgsk 197

RESULT 8  
 W30044  
 ID W30044 standard; Protein; 248 AA.  
 AC W30044;

DT 20-APR-1998 (first entry)

DE Human consensus bikunin.

Human; consensus bikunin; inhibition; trypsin; kallikrein;  
 plasmin; factor XIIa; treatment; prevention; oedema;  
 inflammation; infection; granulomatosis; multiple sclerosis;  
 ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 blood coagulation disease; polytrauma; stroke; haemorrhage;  
 gastric cancer; cervical cancer; metastasis; blood loss.

OS Homo sapiens.  
 XX  
 PN WO9733996-A2.  
 XX  
 PD 18-SEP-1997.  
 XX  
 10-MAR-1997; 97WO-US03894.  
 04-OCT-1996; 96US-0725251.  
 11-MAR-1996; 96US-0013106.  
 14-JUN-1996; 96US-0019793.

(FARB ) BAYER CORP.

PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 DR N-PSDB; T90733.  
 N-PSDB; T90733.

PT New human placental bikunin - used to inhibit kallikrein, trypsin  
 PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 PT perioperative blood loss

PS Claim 1; Page 66; 110pp; English.

XX The present sequence is a consensus human bikunin, which  
CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
CC Bikunin can be used to treat or prevent brain and spinal cord  
CC oedema, inflammation, infection or granulomatosis, multiple  
CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
CC cerebral or subarachnoid haemorrhage and gastric or cervical  
CC cancer and prevent metastasis. It is particularly useful for  
CC reducing blood loss during surgery, and can also be used to treat  
CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
CC influenza and similar viral infections, acute pancreatitis and  
CC gout, and prevent pre-term labour. It has similar properties to  
CC aprotinin, but is less highly charged so should be less  
CC immunogenic and less likely to damage the kidneys. Manipulation  
CC of the bikunin sequence may allow the inhibitory profile to be  
CC altered. It also reduces or eliminates the need for whole donor  
CC blood or blood products during surgery, thereby reducing the risk  
CC of infection and other adverse side effects, as well as reducing  
CC the cost of surgery.

Sequence 248 AA:

Query Match 100.0%; Score 948; DB 18; Length 248;  
Best Local Similarity 100.0%; Pred. No. 1.9e-89; Indels 0; Gaps 0;  
Matches 170; Conservative 0; Mismatches 0;

QY 1 ADRERSIHDFCLVSKVYGRGRASMPRMWYNTVDSQCLFYVGGDGNMNYLTKEECLK 60  
DB 24 adrehsi d fclvskvgrgrasmp rwmw yntv dscqlfyvggdgns mnyltkeekl k 83

QY 61 CATVTENATGDLATSRNAADSSVPSAPRRODSEHSSDMFYERYCTANATGPCRASFP 120  
DB 84 catvt enatgdlatsrnaadssvpsaprr o dsehssdmfy eryct anatgpcrasfp 143

QY 121 RWYFDVERNSCNMFYIGGCRGNKNSYRSEACMLRCFRQENPPLPGSK 170  
DB 144 rwyfdverns c n mfyiggcrgnkn syrseacm lrcfrq enpplpgsk 193

RESULT 9  
W30040 ID W30040 standard; Protein; 252 AA.

AC W30040;

XX 20-APR-1998 (first entry)

Human placental bikunin.

XX Human; placental bikunin; inhibition; trypsin; kallikrein;  
KW plasmin; factor XIIa; treatment; prevention; oedema;  
KW inflammation; infection; granulomatosis; multiple sclerosis;  
KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
KW gastric cancer; cervical cancer; metastasis; blood loss.

OS Homo sapiens.

PN W09733996-A2.

XX 18-SEP-1997.

PF 10-MAR-1997; 97WO-US03894.

PR 04-OCT-1996; 96US-0725251.

PR 11-MAR-1996; 96US-0013106.

PR 14-JUN-1996; 96US-0019793.

XX (FARB) BAYER CORP.

PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;

DR WPI: 1997-470876/43.

DR N-FSDB; T90731.

XX New human placental bikunin - used to inhibit kallikrein, trypsin  
PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
PT perioperative blood loss

PS Claim 1; Page 65; 110pp; English.

XX The present sequence is a human placental bikunin, which  
CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
CC Bikunin can be used to treat or prevent brain and spinal cord  
CC oedema, inflammation, infection or granulomatosis, multiple  
CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
CC cerebral or subarachnoid haemorrhage and gastric or cervical  
CC cancer and prevent metastasis. It is particularly useful for  
CC reducing blood loss during surgery, and can also be used to treat  
CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
CC influenza and similar viral infections, acute pancreatitis and  
CC gout, and prevent pre-term labour. It has similar properties to  
CC aprotinin, but is less highly charged so should be less  
CC immunogenic and less likely to damage the kidneys. Manipulation  
CC of the bikunin sequence may allow the inhibitory profile to be  
CC altered. It also reduces or eliminates the need for whole donor  
CC blood or blood products during surgery, thereby reducing the risk  
CC of infection and other adverse side effects, as well as reducing  
CC the cost of surgery.

Sequence 252 AA:

Query Match 100.0%; Score 948; DB 18; Length 252;  
Best Local Similarity 100.0%; Pred. No. 1.9e-89; Indels 0; Gaps 0;  
Matches 170; Conservative 0; Mismatches 0;

QY 1 ADRERSIHDFCLVSKVYGRGRASMPRMWYNTVDSQCLFYVGGDGNMNYLTKEECLK 60  
DB 28 adrehsi d fclvskvgrgrasmp rwmw yntv dscqlfyvggdgns mnyltkeekl k 87

QY 61 CATVTENATGDLATSRNAADSSVPSAPRRODSEHSSDMFYERYCTANATGPCRASFP 120  
DB 88 catvt enatgdlatsrnaadssvpsaprr o dsehssdmfy eryct anatgpcrasfp 147

QY 121 RWYFDVERNSCNMFYIGGCRGNKNSYRSEACMLRCFRQENPPLPGSK 170  
DB 148 rwyfdverns c n mfyiggcrgnkn syrseacm lrcfrq enpplpgsk 197

RESULT 10  
W13665 ID W13665 standard; Protein; 252 AA.

AC W13665;

XX 11-NOV-1997 (first entry)

DE Hepatocyte growth factor activator inhibitor HAI-II.

XX Hepatocyte growth factor activator inhibitor; HAI-II; HGF; human;

XX protease inhibitor.

OS Homo sapiens.

PF Key Location/Qualifiers

FT Peptide 1..27

FT Protein /label= Sig\_peptide

PN EP758682-A2.

XX 19-FEB-1997.  
 PD  
 XX 23-JUL-1996; 96EP-0111861.  
 PF  
 XX 24-JUL-1995; 95JP-0187134.  
 PR  
 XX (MITU ) MITSUBISHI CHEM CORP.  
 PA  
 XX Kawauchi T, Kitamura N, Shimomura T;  
 PI WPI: 1997-134770/13.  
 DR N-PSDB; T61439.  
 D  
 XX Novel protein HAI-II - inhibits protease activity of hepatocyte  
 PT growth factor activator  
 PS  
 XX Claim 4; Page 18-19; 24pp; English.

This sequence comprises a novel protein, designated HAI-II, that inhibits the protease activity of hepatocyte growth factor (HGF) activator. The sequence was deduced from a cDNA clone (T61439) obtained from cancer cell line MN45. Also claimed are isolated peptides (W13662-64) of HAI-II, the DNA encoding HAI-II, a vector carrying this DNA, and a host cell, pref. an animal cell, transformed with the vector. HAI-II can be used for regulating HGF activator activity (and thus HGF activity) in vitro and in vivo. It may also be used for investigating the function of HAI-II in vivo and the effect of HAI-II in hepatic disorders.

Sequence 252 AA;

Query Match 100.0%; Score 948; DB 19; Length 252;  
 Best Local Similarity 100.0%; Pred. No. 1.9e-89;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADDRESSHDFCLVSKVYVGRASMPRMWYNTDSCQLFVYGGCDGNSNNYLTKEECLK 60  
 Db addresshdfclvskvgrasmprrwvnyvldgscqlfvyggcdgnsnnyltkeecik 87  
 QY 61 CATVTENATGDLATSRNAADSSVPSPARRRODSEHSSDMFNEYECTANAVTGCRASFP 120  
 Db catvtenatgdlatsrnaadssvpaprrqdsehdssdmfneyectanavtgcrafp 147  
 QY 121 RWFYDVERNSCNNFIYGGCGRGNKNSYSEACMLRCRQENPPLPGSK 170  
 Db rwyfdvernschnfiyggcgrgnknsyseacmlrcrfqgenpplpgsk 197

#### RESULT 11

W70286  
 ID W70286 standard; Protein; 252 AA.

AC W70286;

DT 06-NOV-1998 (first entry)

DE Human tissue factor pathway inhibitor-3 (TFPI-3).

XX Human tissue factor pathway inhibitor-3; TFPI-3; blood clot; sepsis;  
 KW fibrin clot; coronary occlusion; acute myocardial infarction;  
 KW prophylaxis; peripheral arterial embolism; inflammatory disease;  
 KW transplant rejection; anticoagulant; blood transfusion;  
 KW extracorporeal circulation; dialysis; haemophili; Kunitz type domain.  
 XX Homo sapiens.

Key Location/Qualifiers  
 FT Peptide 1..27  
 FT /note= "Signal peptide"  
 FT Protein 28..252  
 FT /note= "TFPI-3"

XX WO9833920-A2.  
 PN  
 XX 06-AUG-1998.  
 PD  
 XX 27-JAN-1998; 98WO-US01468.  
 PF  
 XX 31-JAN-1997; 97US-0036703.  
 PR  
 XX (HUMA-) HUMAN GENOME SCI INC.  
 PA  
 XX Gentz RL, Hsu T, Ni J, Rosen CA;  
 PI WPI: 1998-437473/37.  
 DR N-PSDB; V33063.  
 D  
 XX Isolated tissue factor pathway inhibitor-3 - used to develop  
 PT products for treating, e.g. pulmonary embolism, thrombosis, sepsis,  
 PT inflammatory disease, transplant rejection or haemophilia  
 PS Disclosure; Fig 1A-1B; 57pp; English.

The present sequence represents a human tissue factor pathway inhibitor-3 (TFPI-3) which contains two Kunitz type domains. The invention also provides the TFPI-3 cDNA and screening methods for identifying agonists and antagonists of TFPI-3. As TFPI-3 inhibits protease activity, it is claimed to be useful for, e.g. inhibiting intravascular clotting and preventing the formation of fibrin clots both in vitro and in vivo, for treating coronary occlusion with acute myocardial infarction and in the prophylaxis and treatment of diseases and transplant rejection. TFPI-3 is also claimed to be useful as an anticoagulant in blood transfusions, extracorporeal circulation, and dialysis procedures and in blood samples for laboratory purposes. The TFPI-3 antagonists are claimed to be useful for promoting coagulation, e.g. in the treatment of haemophilia.

Sequence 252 AA;

Query Match 100.0%; Score 948; DB 19; Length 252;  
 Best Local Similarity 100.0%; Pred. No. 1.9e-89;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADDRESSHDFCLVSKVYVGRASMPRMWYNTDSCQLFVYGGCDGNSNNYLTKEECLK 60  
 Db addresshdfclvskvgrasmprrwvnyvldgscqlfvyggcdgnsnnyltkeecik 87  
 QY 61 CATVTENATGDLATSRNAADSSVPSPARRRODSEHSSDMFNEYECTANAVTGCRASFP 120  
 Db catvtenatgdlatsrnaadssvpaprrqdsehdssdmfneyectanavtgcrafp 147  
 QY 121 RWFYDVERNSCNNFIYGGCGRGNKNSYSEACMLRCRQENPPLPGSK 170  
 Db rwyfdvernschnfiyggcgrgnknsyseacmlrcrfqgenpplpgsk 197

#### RESULT 12

W30051  
 ID W30051 standard; Protein; 153 AA.

AC W30051;

DT 20-APR-1998 (first entry)

DE Human placental bikunin.

XX Human; placental bikunin; inhibition; trypsin; kallikrein;  
 KW plasmin; factor XIII; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.

XX Homo sapiens.  
 OS WO9733996-A2.  
 XX 18-SEP-1997.  
 XX 10-MAR-1997: 97WO-US03894.  
 XX 04-OCT-1996: 96US-0725251.  
 XX 11-MAR-1996: 96US-0013106.  
 XX 14-JUN-1996: 96US-0019793.  
 XX (FARB ) BAYER CORP.  
 XX Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 PI WPI; 1997-470876/43.  
 XX New human placental bikunin - used to inhibit kallikrein, trypsin  
 etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 perioperative blood loss  
 Claim 1; Page 67; 110pp; English.  
 PS The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor Xlla.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.  
 CC Sequence 153 AA;  
 SO

Query Match 90.6%; Score 859; DB 18; Length 153;  
 Best Local Similarity 100.0%; Pred. No. 1.3e-80;  
 Matches 153; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 IHDFCLVSKVYGRGRASMPRMWYNVTGSCQLFVYGGCGNSNNYLTKECLKCAVTE 66  
 |||||||  
 Db 1 ihdfclvskvgrcraemprrwvnyvtdgscqlfvygdcgnsnyltkeclkkcatve 60  
 QY 67 NATGDLATSRNADSVPSAPRRODSEHSDMFNEEYETANAVYGPCRASFPWYDFV 126  
 |||||||  
 Db 61 natgdlatsrnaadsvpsaprrdgsedhsdmfneyeeyctanavtgcrafpwydfv 120  
 QY 127 ERNSCNFIYGGCRGNKNSYRSEACMLRCFRQ 159  
 |||||||  
 Db 121 ernscnfiyggcrgnknsyrseacmlrcfrq 153

RESULT 13  
 W30052  
 ID W30052 standard; Protein; 146 AA.  
 XX W30052;  
 AC W30052;  
 XX 20-APR-1998 (first entry)  
 DT  
 XX

DE Human placental bikunin.  
 XX Human; placental bikunin; inhibition; trypsin; kallikrein;  
 KW plasmin; factor Xlla; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.  
 XX Homo sapiens.  
 OS WO9733996-A2.  
 XX 18-SEP-1997.  
 XX 10-MAR-1997: 97WO-US03894.  
 XX 04-OCT-1996: 96US-0725251.  
 XX 11-MAR-1996: 96US-0013106.  
 XX 14-JUN-1996: 96US-0019793.  
 XX (FARB ) BAYER CORP.  
 XX Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 PI WPI; 1997-470876/43.  
 XX New human placental bikunin - used to inhibit kallikrein, trypsin  
 etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 perioperative blood loss  
 Claim 1; Page 67; 110pp; English.  
 PS The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor Xlla.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.  
 CC Sequence 146 AA;  
 SO

Query Match 86.4%; Score 819; DB 18; Length 146;  
 Best Local Similarity 100.0%; Pred. No. 1.6e-76;  
 Matches 146; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 CLVSKVYGRGRASMPRMWYNVTGSCQLFVYGGCGNSNNYLTKECLKCAVTEATG 70  
 |||||||  
 Db 1 clvskvgrcraemprrwvnyvtdgscqlfvygdcgnsnyltkeclkkcatvtenatg 62  
 QY 71 DLATSRNADSVPSAPRRODSEHSDMFNEEYETANAVYGPCRASFPWYDFVDRNS 130  
 |||||||  
 Db 61 dlatsrnaadsvpsaprrdgsedhsdmfneyeeyctanavtgcrafpwydfvdrns 120  
 QY 131 CNMFIYGGCRGNKNSYRSEACMLRC 156  
 |||||||  
 Db 121 cnmfiyggcrgnknsyrseacmlrc 146

RESULT	14
ID	W30061
XX	
XX	W30061 standard; Protein; 170 AA.
AC	
DT	W30061;
XX	
XX	20-APR-1998 (first entry)
DE	
XX	Human consensus bikunin.
KW	
KW	Human; consensus bikunin; inhibition; trypsin; kallikrein;
KW	plasma; factor XIa; treatment; prevention; oedema;
KW	inflammation; infection; granulomatosis; multiple sclerosis;
KW	ischemia; perioperative blood loss; sepsis; shock; fibrinolysis;
KW	blood coagulation disease; polytrauma; stroke; haemorrhage;
KW	gastric cancer; cervical cancer; metastasis; blood loss.
OS	
XX	Homo sapiens.
XX	
Key	Location/Qualifiers
Misc-difference	1..170
/note=	"Xaa = any amino acid, except Cys provided that at least one is different from the corresponding native amino acid"
WT	
FT	
FT	
FT	
XX	
PX	MO9733996-A2.
PD	
PD	18-SEP-1997.
PE	
PE	10-MAR-1997; 97WO-USG03894.
PR	
PR	04-OCT-1996; 96US-0725251.
PR	11-MAR-1996; 96US-0013106.
PR	14-JUN-1996; 96US-0019793.
PA	(PARB) BAYER CORP.
XX	
PI	Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;
DR	
XX	WPI; 1997-470876/43.
PT	
PT	New human placental bikunin - used to inhibit kallikrein, trypsin
PT	etc. in treatment of oedema, multiple sclerosis, fibrosis, or
XX	perioperative blood loss
XX	
PS	
XX	Disclosure; Page 10; 110pp; English.
XX	
XX	The present sequence is a consensus human bikunin, which
XX	inhibits, e.g. trypsin, kallikrein, plasma and factor XIa.
CC	Bikunin can be used to treat or prevent brain and spinal cord
CC	oedema, inflammation, infection or granulomatosis, multiple
CC	sclerosis, ischaemia, perioperative blood loss, sepsis, shock,
CC	fibrosis, blood coagulation diseases, polytrauma, stroke,
CC	cerebral or subarachnoid haemorrhage and gastric or cervical
CC	cancer and prevent metastasis. It is particularly useful for
CC	reducing blood loss during surgery, and can also be used to treat
CC	other cancer, arthritis, anaemia, non-insulin dependent diabetes,
CC	gout, and prevent pre-term labour. It has similar properties to
CC	aprotinin, but is less highly charged so should be less
CC	immunogenic and less likely to damage the kidneys. Manipulation
CC	of the bikunin sequence may allow the inhibitory profile to be
CC	altered. It also reduces or eliminates the need for whole donor
CC	blood or blood products during surgery, thereby reducing the risk
CC	of infection and other adverse side effects, as well as reducing
CC	the cost of surgery.
XX	
Sequence	170 AA;
XX	

Query Match	79.1%	Score 750;	DB 18;	Length 170;
Best Local Similarity	81.2%;	Pred. No. 2.2e69;		
Matches 138;	Conservative 0;	Mismatches 32;	Indels 0;	Gaps 0

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Db	1	adterstlxdiclyskvxcgccccccccwwwjyvtidgscqlfxyxgcxxxxsmnyxtkeecLk	60
QY	61	CAVTEVATGDGLTSRRADSSVPSAPRODSDSDHSDMFNVEYCYTANNAVTPCRAFP	120
Db	61	catxtenatgldltsrmaadssvpsaprrqdsedhsdntfnyxeyctanaavxgccccxxx	120
QY	121	RMTEFVERNSCNFFYGGCGNGKNSYRSEACMLRCFROENPPLPLGSK	170
Db	121	xwffdvcrnscafnfxyxgcxxxknsyxseacmlrfrfxqenpplpLgsk	170
RESULT	15		
ID	W30054		
XX	W30054 standard; Protein: 92 AA.		
AC	W30054:		
XX			
XX	20-APR-1998 (first entry)		
DE			
XX	Human placental bikunin.		
KW	Human; placental bikunin; inhibition; trypsin; kallikrein;		
KW	plasmaIn; factor XIIa; treatment; prevention; oedema;		
KW	inflammation; infection; granulomatosis; multiple sclerosis;		
KW	ischæmia; perioperative blood loss; sepsis; shock; fibrosis;		
KW	blood coagulation disease; polycytræmia; stroke; hæmorrhage;		
KW	gastric cancer; cervical cancer; metastasis; blood loss.		
OS	Homo sapiens.		
PN			
XX	W09733996-A2.		
XX			
PD	18-SEP-1997.		
XX			
PF	10-MAR-1997; 97MO-US03894.		
XX			
PR	04-OCT-1996; 96GUS-0725251.		
PR	11-MAR-1996; 96GUS-0013106.		
PR	14-JUN-1996; 96GUS-0019793.		
XX			
PA	(FARB ) BAYER CORP.		
PI			
PI	Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;		
DR	WPI; 1997-470876/43.		
XX			
PT			
PT	New human placental bikunin - used to inhibit kallikrein, trypsin		
PT	etc. in treatment of oedema, multiple sclerosis, fibrosis, or		
PS	perioperative blood loss		
XX			
XX	Claim 1: Page 67; 110pp; English.		
CC	The present sequence is a human placental bikunin, which		
CC	inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.		
CC	Bikunin can be used to treat or prevent brain and spinal cord		
CC	oedema, inflammation, infection or granulomatosis, multiple		
CC	fibrosis, ischæmia, perioperative blood loss, sepsis, shock,		
CC	cerebral or subarachnoid hæmorrhage and gastric or cervical		
CC	cancer and prevent metastasis. It is particularly useful for		
CC	reducing blood loss during surgery, and can also be used to treat		
CC	other cancer, arthritis, anaemia, non-insulin dependent diabetes,		
CC	influenza and similar viral infections, acute pancreatitis and		
CC	gout, and prevent pre-term labour. It has similar properties to		
CC	aprotinin, but is less highly charged so should be less		
CC	immunogenic and less likely to damage the kidneys. Manipulation		
CC	of the bikunin sequence may allow the inhibitory profile to be		
CC	altered. It also reduces or eliminates the need for whole donor		
CC	blood or blood products during surgery, thereby reducing the risk		
CC	of infection and other adverse side effects.		

RESULT 15  
 ID W30054  
 AC W30054 standard; Protein: 92 AA.  
 XX W30054;  
 DI 20-APR-1998 (first entry)  
 DE Human placental bikunin.  
 KW Human; placental bikunin; inhibition; trypsin; kallikrein;  
 KW plasmin; factor Xlla; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; hemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.  
 XX  
 OS Homo sapiens.  
 PN W09733996-A2.  
 XX  
 PD 18-SEP-1997.  
 XX  
 PF 10-MAR-1997; 97MO-US03894.  
 XX  
 PR 04-OCT-1996; 96US-0725251.  
 PR 11-MAR-1996; 96US-0013106.  
 PR 14-JUN-1996; 96US-0019793.  
 XX  
 PA (FARB ) BAYER CORP.  
 XX  
 PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PJ.  
 XX  
 DR WPI; 1997-470876/43.  
 XX  
 PT New human placental bikunin - used to inhibit kallikrein, trypsin  
 PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 PT perioperative blood loss  
 XX  
 PS Claim 1; Page 67; 110pp; English.  
 XX  
 CC The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor Xlla.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing





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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:03:28 ; Search time 12.37 seconds

(without alignments)  
246,782 Million cell updates/sec

Title: US-09-441-654A-1

Sequence: 1 ADRESDHDFCLVSKVVGRC.....ACMDRCFROQENPLPLGSK 170

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 174772 seqs, 17957048 residues

Total number of hits satisfying chosen parameters: 174772

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued\_Patents\_AA:\*  
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3: /cgn2\_6/ptodata/2/1aa/6.COMB.pep:\*  
4: /cgn2\_6/ptodata/2/1aa/PTCUTS.COMB.pep:\*  
5: /cgn2\_6/ptodata/2/1aa/Backfile1.pep:\*

Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	948	100.0	252	1	US-08-685-660A-7
2	948	100.0	252	1	US-08-974-196-7
3	247.5	26.1	122	2	US-08-422-333-12
4	247.5	26.1	122	5	5187153-20
5	247.5	26.1	122	5	5220013-23
6	244.5	25.8	143	2	US-08-422-333-10
7	244.5	25.8	143	5	5223482-20
8	244.5	25.8	144	5	5187153-18
9	244.5	25.8	147	1	US-08-358-160-72
10	243.5	25.7	127	5	5466783-24
11	241.5	25.5	123	5	5466783-21
12	233.5	24.6	122	5	5223482-22
13	233.5	24.6	276	1	US-07-828-920A-1
14	233.5	24.6	276	1	US-08-437-841-9
15	233.5	24.6	276	1	US-08-286-521-9
16	233.5	24.6	276	1	US-08-436-175-9
17	233.5	24.6	276	2	US-08-796-850-1
18	233.5	24.6	276	2	US-08-436-175-9
19	233.5	24.6	276	3	US-08-854-764-3
20	233.5	24.6	276	4	PCR-US95-09377-3
21	233.5	24.6	277	4	US-07-844-297-1
22	233.5	24.6	304	1	US-08-026-145-2
23	233.5	24.6	304	1	US-08-446-646-9
24	233.5	24.6	304	1	US-08-676-125A-18
25	233.5	24.6	304	2	US-09-136-012A-18
26	233.5	24.6	304	2	US-08-676-124-1
27	233.5	24.6	304	3	US-08-208-264A-25
28	233.5	24.6	304	3	US-09-414-878-1

29	233.5	24.6	304	3	US-09-240-136-1	Sequence 1, Appl
30	233.5	24.6	304	5	5466783-2	Patent No. 5466783
31	233.5	24.6	352	3	US-08-854-764-2	Sequence 2, Appl
32	233.5	24.6	352	4	PCR-US95-09377-2	Sequence 2, Appl
33	232.5	24.5	123	5	5466783-22	Patent No. 5466783
34	232	24.5	161	1	US-08-437-841-19	Sequence 19, Appl
35	232	24.5	161	1	US-08-286-521-19	Sequence 19, Appl
36	232	24.5	161	1	US-08-436-175-19	Sequence 19, Appl
37	232	24.5	161	4	PCR-US95-09464-19	Sequence 19, Appl
38	230	24.3	122	5	5466783-23	Patent No. 5466783
39	228	24.1	183	1	US-07-828-920A-5	Sequence 1, Appl
40	226	23.8	189	1	US-07-828-920A-7	Sequence 1, Appl
41	219	23.1	213	5	5466783-25	Patent No. 5466783
42	214.5	22.6	213	2	US-08-796-850-2	Sequence 2, Appl
43	214.5	22.6	235	1	US-08-147-710-2	Sequence 2, Appl
44	214.5	22.6	235	1	US-08-458-090-2	Sequence 2, Appl
45	214.5	22.6	235	2	US-08-457-887-2	Sequence 2, Appl

## ALIGNMENTS

RESULT 1  
US-08-685-660A-7  
; Sequence 7, Application US/08685660A  
; Patent No. 5731412  
; GENERAL INFORMATION:  
; APPLICANT: SHIMOMURA, Takeshi  
; APPLICANT: KAWAGUCHI, Toshiya  
; APPLICANT: KITAMURA, Naomi  
; TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
; TITLE OF INVENTION: AND METHOD OF PRODUCING THE PROTEIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: SOGHROE, MION, ZINN, MACPEAK & SEAS  
; STREET: 2100 Pennsylvania Avenue, N.W.  
; CITY: Washington  
; STATE: DC  
; COUNTRY: USA  
; ZIP: 20037  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy Disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patentin Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/685,660A  
; FILING DATE: 24-JUL-1996  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: JPA Hel 7-187134  
; FILING DATE: 24-JUL-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: KIT, Gordon  
; REGISTRATION NUMBER: 30,764  
; REFERENCE/DOCKET NUMBER: Q-42295  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (202) 293-7060  
; TELEFAX: (202) 293-7860  
; INFORMATION FOR SEQ ID NO: 7:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 252 amino acids  
; TYPE: amino acid  
; TOPOLOGY: linear  
; MOLECULE TYPE: peptide  
; US-08-685-660A-7

Query Match 100.0%; Score 948; DB 1; Length 252;  
Best Local Similarity 100.0%; Pred. No. 3e-92;

Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 ADRESDHDFCLVSKVVGRCASPMRWNYNTDSCGFYGGCGGNSNNTLTBECLAK 60  
|||||

Wed Jan 31 15:19:33 2001

us-09-441-654a-1.ra1

Page 2

Db 28 ADERSIHDFCLVSKVYGRCRASMPRWYNTVDGSCQLFYGGCGDGSNNYLKKECLK 87  
QY 61 CATVTENATGDLATSRNADSSVPSAPRRDSEDHSSDMFNEYECTANAVTGPCRASFP 120  
Db 88 CATVTENATGDLATSRNADSSVPSAPRRDSEDHSSDMFNEYECTANAVTGPCRASFP 147  
QY 121 RWYFDVERNSCNFFIYGGCGRGNKNSYRSEACMLRCFROENPPLPLGSK 170  
Db 148 RWYFDVERNSCNFFIYGGCGRGNKNSYRSEACMLRCFROENPPLPLGSK 197  
RESULT 2  
US-08-974-196-7  
; Sequence 7, Application US/08974196  
; Patent No. 5854396  
; GENERAL INFORMATION:  
; APPLICANT: SHIMOMURA, Takeshi  
; APPLICANT: KAWAGUCHI, Toshiya  
; APPLICANT: KITAMURA, Naomi  
; TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR "SAME  
; TITLE OF INVENTION: AND METHOD OF PRODUCING THE PROTEIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS  
; STREET: 2100 Pennsylvania Avenue, N.W.  
; CITY: Washington  
; STATE: DC  
; COUNTRY: USA  
; ZIP: 20037  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/974,196  
; FILING DATE:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/685,660  
; FILING DATE: 24-JUL-1996  
; APPLICATION NUMBER: JPA Hei 7-187134  
; FILING DATE: 24-JUL-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: KIT, Gordon  
; REGISTRATION NUMBER: 30,764  
; REFERENCE/DOCKET NUMBER: Q-42295  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (202) 293-7060  
; TELEFAX: (202) 293-7860  
; INFORMATION FOR SEQ ID NO: 7:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 252 amino acids  
; TYPE: amino acid  
; TOPOLOGY: linear  
; MOLECULE TYPE: peptide  
US-08-974-196-7  
Query Match 100.0%; Score 948; DB 2; Length 252;  
Best Local Similarity 100.0%; Pred. No. 3e-92; Indels 0; Gaps 0;  
Matches 170; Conservative 0; Mismatches 0;  
QY 1 ADERSIHDFCLVSKVYGRCRASMPRWYNTVDGSCQLFYGGCGDGSNNYLKKECLK 60  
Db 28 ADERSIHDFCLVSKVYGRCRASMPRWYNTVDGSCQLFYGGCGDGSNNYLKKECLK 87  
QY 61 CATVTENATGDLATSRNADSSVPSAPRRDSEDHSSDMFNEYECTANAVTGPCRASFP 120  
Db 88 CATVTENATGDLATSRNADSSVPSAPRRDSEDHSSDMFNEYECTANAVTGPCRASFP 147  
QY 121 RWYFDVERNSCNFFIYGGCGRGNKNSYRSEACMLRCFROENPPLPLGSK 170  
Db 148 RWYFDVERNSCNFFIYGGCGRGNKNSYRSEACMLRCFROENPPLPLGSK 197

Db 3  
RESULT 3  
US-08-422-333-12  
; Sequence 12, Application US/08422333  
; Patent No. 5912410  
; GENERAL INFORMATION:  
; APPLICANT: CORDELL, Barbara L.  
; TITLE OF INVENTION: TRANSGENIC NON-HUMAN MAMMAL DISPLAYING  
; TITLE OF INVENTION: THE ANTIOID-FORMING PATHOLOGY OF ALZHEIMER'S DISEASE  
; NUMBER OF SEQUENCES: 30  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Scios, Inc.  
; STREET: 2450 Bayshore Parkway  
; CITY: Mountain View  
; STATE: CA  
; COUNTRY: USA  
; ZIP: 94043  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/422,333  
; FILING DATE: 13-APR-1995  
; CLASSIFICATION: 800  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Shearer, Peter R.  
; REGISTRATION NUMBER: 28,117  
; REFERENCE/DOCKET NUMBER: 21900-28048.00  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 966-1550  
; TELEFAX: (415) 968-2438  
; INFORMATION FOR SEQ ID NO: 12:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 122 amino acids  
; TYPE: amino acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
US-08-422-333-12  
Query Match 26.1%; Score 247.5; DB 2; Length 122;  
Best Local Similarity 32.4%; Pred. No. 6e-19;  
Matches 48; Conservative 16; Mismatches 45; Indels 39; Gaps 1;  
QY 9 DFLVSKVYGRCRASMPRWYNTVDGSCQLFYGGCGDGSNNYLKKECLKCATYTENA 68  
Db 3 DSCQLDYSGPCLGLFKRYFNGTSMACETFLYGGCGMGLNLFLOKBELOCTRTV----- 58  
QY 69 TGDPLATSRNADSSVPSAPRRDSEDHSSDMFNEYECTANAVTGPCRASFPRWYFDVER 128  
Db 59 -----EACNLPLIVGCPICRAFIOLMADAVK 83  
QY 129 NSCNFFIYGGCGRGNKNSYRSEACMLRC 156  
Db 84 GKCYRFSYGGCGRGNKNSYRSEACMLRC 111  
RESULT 4  
5187153-20  
; Patent No. 5187153  
; APPLICANT: CORDELL, BARBARA, SCHILLING, JAMES W., KITAMURA, NOBUHIKO  
; TITLE OF INVENTION: METHODS OF TREATMENT USING ALZHEIMER'S  
; ANTIOID POLYPEPTIDE DERIVATIVES  
; NUMBER OF SEQUENCES: 33  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/07/502,273  
; FILING DATE: 29-MAR-1990  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 361,912  
; FILING DATE: 06-JUN-1989

APPLICATION NUMBER: 359,911  
 FILING DATE: 12-MAY-1989  
 APPLICATION NUMBER: 87,002  
 FILING DATE: 18-AUG-1987  
 APPLICATION NUMBER: 8,810  
 FILING DATE: 30-JAN-1987  
 APPLICATION NUMBER: 948,376  
 FILING DATE: 31-DEC-1986  
 APPLICATION NUMBER: 932,193  
 FILING DATE: 17-NOV-1986  
 SEQ ID NO: 20:  
 LENGTH: 122  
 5187153-20

Query Match 26.1%; Score 247.5; DB 5; Length 122;  
 Best Local Similarity 32.4%; Pred. No. 6e-19; Indels 39; Gaps 1;  
 Matches 48; Conservative 16; Mismatches 45;

9 DECLVSKVYGRCRASMPRWYNTDSCOLFVYGGCDGNSNNYLTKEECLKCATYTENA 68  
 3 DSCOLDYSGQPCGLGFKRYFNGTSMACETFLYGGCMGNINFLSQKECLQTCRTV---- 58  
 69 TGDILATSRNADSSVSPAPRRDSEHSSDMFNEYECTANAVTGRCRASFPWYFVER 128  
 59 -----EACNLPYVGPCRAFIQIMAFDAVK 83  
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 84 GKCVRFSGGCKGNKNGKFSQKECKEYC 111

RESULT 5  
 5220013-23  
 Patent No. 5220013  
 APPLICANT: PONTE, PHYLLIS A.;CORDELL, BARBARA  
 TITLE OF INVENTION: DNA SEQUENCE USEFUL FOR THE DETECTION  
 OF ALZHEIMER'S DISEASE  
 NUMBER OF SEQUENCES: 30  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/07/444,118  
 FILING DATE: 30-NOV-1989  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: 87,002  
 FILING DATE: 18-AUG-1987  
 APPLICATION NUMBER: 8,810  
 FILING DATE: 30-JAN-1987  
 APPLICATION NUMBER: 948,376  
 FILING DATE: 31-DEC-1986  
 APPLICATION NUMBER: 932,193  
 FILING DATE: 17-NOV-1986  
 SEQ ID NO: 23:  
 LENGTH: 122  
 5220013-23

Query Match 26.1%; Score 247.5; DB 5; Length 122;  
 Best Local Similarity 32.4%; Pred. No. 6e-19; Indels 39; Gaps 1;  
 Matches 48; Conservative 16; Mismatches 45;

9 DECLVSKVYGRCRASMPRWYNTDSCOLFVYGGCDGNSNNYLTKEECLKCATYTENA 68  
 3 DSCOLDYSGQPCGLGFKRYFNGTSMACETFLYGGCMGNINFLSQKECLQTCRTV---- 58  
 69 TGDILATSRNADSSVSPAPRRDSEHSSDMFNEYECTANAVTGRCRASFPWYFVER 128  
 59 -----EACNLPYVGPCRAFIQIMAFDAVK 83  
 129 NSCNNFIYGGCRGNKNSYSEACMLRC 156  
 84 GKCVRFSGGCKGNKNGKFSQKECKEYC 111

RESULT 6  
 US-08-422-333-10  
 Sequence 10, Application US/08422333  
 Patent No. 5912410  
 GENERAL INFORMATION:  
 APPLICANT: CORDELL, Barbara L.  
 TITLE OF INVENTION: TRANSGENIC NON-HUMAN MAMMAL DISPLAYING  
 THE AMYLOID-FORMING PATHOLOGY OF ALZHEIMER'S DISEASE  
 NUMBER OF SEQUENCES: 30  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Scios, Inc.  
 STREET: 2450 Bayside Parkway  
 CITY: Mountain View  
 STATE: CA  
 COUNTRY: USA  
 ZIP: 94043  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 SOFTWARE: Patent Release #1.0, Version #1.30  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/422,333  
 FILING DATE: 13-APR-1995  
 CLASSIFICATION: 800  
 ATTORNEY/AGENT INFORMATION:  
 NAME: Shearer, Peter R.  
 REGISTRATION NUMBER: 28,117  
 REFERENCE/DOCKET NUMBER: 21900-28048.00  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: (415) 966-1550  
 TELEFAX: (415) 968-2438  
 INFORMATION FOR SEQ ID NO: 10:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 143 amino acids  
 TYPE: amino acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 US-08-422-333-10

Query Match 25.8%; Score 244.5; DB 2; Length 143;  
 Best Local Similarity 31.8%; Pred. NO. 1.5e-18;  
 Matches 47; Conservative 15; Mismatches 47; Indels 39; Gaps

9 DECLVSKVYGRCRASMPRWYNTDSCOLFVYGGCDGNSNNYLTKEECLKCATYTENA 68  
 24 DSCOLDYSAGCMGNTSRYFNGTSMACETFOYGGCMGNFVTEKECLQTCRTVAA-- 81  
 69 TGDILATSRNADSSVSPAPRRDSEHSSDMFNEYECTANAVTGRCRASFPWYFVER 128  
 82 -----CNLPVIRGCRAFIQLMAFDAYK 104  
 129 NSCNNFIYGGCRGNKNSYSEACMLRC 156  
 105 GKCVLPFYGGCGGNGKNGKFSQKECKEYC 132

RESULT 7  
 5223482-20  
 Patent No. 5223482  
 APPLICANT: SCHILLING, JAMES W.;PONTE, PHYLLIS A.;CORDELL,  
 BARBARA  
 TITLE OF INVENTION: RECOMBINANT ALZHEIMER'S PROTEASE  
 INHIBITOR AMYLOID PROTEIN AND METHOD OF USE  
 NUMBER OF SEQUENCES: 34  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/07/361,912  
 FILING DATE: 06-JUN-1989  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: 359,911  
 FILING DATE: 12-MAY-1989  
 APPLICATION NUMBER: 87,002

;; FILING DATE: 18-AUG-1987  
;; APPLICATION NUMBER: 8,810  
;; FILING DATE: 30-JAN-1987  
;; APPLICATION NUMBER: 948,376  
;; FILING DATE: 31-DEC-1986  
;; APPLICATION NUMBER: 932,193  
;; FILING DATE: 17-NOV-1986  
;; SEQ ID NO: 20  
;; LENGTH: 143  
5223482-20

Query Match  
Best Local Similarity 25.8%; Score 244.5; DB 5; Length 143;  
Matches 47; Conservative 15; Mismatches 47; Indels 39; Gaps 1;

QY 9 DCLVSKVVGRCRASPMPRMWYNTDSCQLFYVGGCDGNSNNYLTKECKLKCATVTENA 68  
DB 24 DSCQIGYSAGPCMGMTSRFYNGTSMACETFOYGGCGMGNNNVTIEKCIQTCTVAA-- 81  
69 TGDLSRNAADSSVPSAPRRDSEHSSDMFYEEYCTANAVTGPCRASFPWMYFDVER 128  
DB 82 -----CNLPYIRGPCRAFIOLMADAVK 104

QY 129 NSCNNFIYGGCRGNKNSYSEACMLRC 156  
DB 105 GKCVLFPGYGGCGNGNKFYSEKCEKREYC 132

RESULT 8  
5187153-18  
PATENT NO. 5187153  
APPLICANT: CORDELL, BARBARA; SCHILLING, JAMES W.; KATUNUMA, NOBUHIKO  
TITLE OF INVENTION: METHODS OF TREATMENT USING ALZHEIMER'S  
AMLODIPOLYMERPEPTIDE DERIVATIVES  
NUMBER OF SEQUENCES: 33  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/502,273  
FILING DATE: 29-MAR-1990  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 361,912  
FILING DATE: 06-JUN-1986  
APPLICATION NUMBER: 359,911  
FILING DATE: 12-MAY-1989  
APPLICATION NUMBER: 87,002  
FILING DATE: 18-AUG-1987  
APPLICATION NUMBER: 8,810  
FILING DATE: 30-JAN-1987  
APPLICATION NUMBER: 948,376  
FILING DATE: 31-DEC-1986  
APPLICATION NUMBER: 932,193  
FILING DATE: 17-NOV-1986  
SEQ ID NO: 18  
LENGTH: 144  
5187153-18

Query Match  
Best Local Similarity 25.8%; Score 244.5; DB 5; Length 144;  
Matches 47; Conservative 15; Mismatches 47; Indels 39; Gaps 1;

QY 9 DCLVSKVVGRCRASPMPRMWYNTDSCQLFYVGGCDGNSNNYLTKECKLKCATVTENA 68  
DB 24 DSCQIGYSAGPCMGMTSRFYNGTSMACETFOYGGCGMGNNNVTIEKCIQTCTVAA-- 81  
69 TGDLSRNAADSSVPSAPRRDSEHSSDMFYEEYCTANAVTGPCRASFPWMYFDVER 128  
DB 82 -----CNLPYIRGPCRAFIOLMADAVK 104  
QY 129 NSCNNFIYGGCRGNKNSYSEACMLRC 156  
DB 105 GKCVLFPGYGGCGNGNKFYSEKCEKREYC 132

RESULT 9  
US-08-358-160-72  
Sequence 72, Application US/08358160  
Patent No. 5663143  
GENERAL INFORMATION:

APPLICANT: LEY, Arthur C.  
APPLICANT: LADNER, Robert C.  
APPLICANT: GUTERMAN, Sonia K.  
APPLICANT: ROBERTS, Bruce L.  
APPLICANT: MARKLAND, William  
APPLICANT: KENT, Rachel B.  
TITLE OF INVENTION: ENGINEERED HUMAN-DERIVED KUNITZ  
NUMBER OF SEQUENCES: 234  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: BROWDY AND NEIMARK  
STREET: 419 Seventh Street, N.W. Suite 300  
CITY: Washington  
STATE: District of Columbia  
COUNTRY: USA  
ZIP: 20004

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/358,160  
FILING DATE: 16-DEC-1994  
CLASSIFICATION: 514  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/133,031  
FILING DATE: 13-OCT-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/009,319  
FILING DATE: 26-JAN-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/664,989  
FILING DATE: 01-MAR-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/487,063  
FILING DATE: 02-MAR-1990  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/240,160  
FILING DATE: 02-SEP-1988  
ATTORNEY/AGENT INFORMATION:  
NAME: Cooper, Iyer P.  
REGISTRATION NUMBER: 28,005  
REFERENCE/DOCKET NUMBER: LEY-1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-628-5197  
TELEFAX: 202-737-3528  
TELEX: 248633

INFORMATION FOR SHD ID NO: 72:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 147 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
US-08-358-160-72

Query Match  
Best Local Similarity 32.4%; Score 244.5; DB 1; Length 147;  
Matches 48; Conservative 14; Mismatches 47; Indels 39; Gaps 1;

QY 9 DCLVSKVVGRCRASPMPRMWYNTDSCQLFYVGGCDGNSNNYLTKECKLKCATVTENA 68  
DB 24 DSCQIGYSAGPCMGMTSRFYNGTSMACETFOYGGCGMGNNNVTIEKCIQTCTVAA-- 81  
69 TGDLSRNAADSSVPSAPRRDSEHSSDMFYEEYCTANAVTGPCRASFPWMYFDVER 128

Db 82 -----CNLPVIRGPCRAFIQLWAFDAVK 104  
QY 129 NSCNFIYGGCGNKNYSRSEACMLRC 156  
Db 105 GKCVLPYGGCGNGNKNFYSEKCEYIC 132  
RESULT 10  
5466783-24  
; Patent No. 5466783  
; APPLICANT: Wu, Tze-Chen.; Kretzmer, Kuniko K.; Broze,  
; George J., Jr.  
; TITLE OF INVENTION: HUMAN TISSUE FACTOR INHIBITOR  
; NUMBER OF SEQUENCES: 26  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/93,285  
; FILING DATE: 15-JUL-1993  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 566,280  
; FILING DATE: 13-AUG-1990  
; APPLICATION NUMBER: 123,753  
; FILING DATE: 23-NOV-1987  
; APPLICATION NUMBER: 77,366  
; FILING DATE: 23-JUL-1987  
; SEQ ID NO: 24:  
; LENGTH: 127  
5466783-24

Query Match 25.7%; Score 243.5; DB 5; Length 127;  
Best Local Similarity 32.7%; Pred. No. 1.7e-18;  
Matches 48; Conservative 14; Mismatches 46; Indels 39; Gaps 1;  
QY 9 DFCLVSKVVGCRASPRWYNTDGSQCLFYVGGCGNSNNYLTKEBCLKKCATVTENA 68  
Db 3 DSCOLGYSAGPCGMGTSRYFYNGTSMACETFYGGCGNGNMFYTEKECLQTCRTVAA-- 60  
QY 69 TGLATSRNAADSVPSAPRRQDSHSDMFNFEYCTANAVTGPCRASFFRWYFDVER 128  
Db 61 -----CNLPVIRGPCRAFIQLWAFDAVK 83

QY 129 NSCNFIYGGCGNKNYSRSEACMLR 155  
Db 84 GKCVLPYGGCGNGNKNFYSEKCECR 110  
RESULT 11  
5466783-21  
; Patent No. 5466783  
; APPLICANT: Wu, Tze-Chen.; Kretzmer, Kuniko K.; Broze,  
; George J., Jr.  
; TITLE OF INVENTION: HUMAN TISSUE FACTOR INHIBITOR  
; NUMBER OF SEQUENCES: 26  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/93,285  
; FILING DATE: 15-JUL-1993  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 566,280  
; FILING DATE: 13-AUG-1990  
; APPLICATION NUMBER: 123,753  
; FILING DATE: 23-NOV-1987  
; APPLICATION NUMBER: 77,366  
; FILING DATE: 23-JUL-1987  
; SEQ ID NO: 21:  
; LENGTH: 123  
5466783-21

Query Match 25.5%; Score 241.5; DB 5; Length 123;  
Best Local Similarity 31.8%; Pred. No. 2.6e-18;  
Matches 47; Conservative 15; Mismatches 47; Indels 39; Gaps 1;

QY 9 DFCLVSKVVGCRASPRWYNTDGSQCLFYVGGCGNSNNYLTKEBCLKKCATVTENA 68  
Db 3 DSCOLGHAQPCGLGMSRYSFYNGTSMACETFYGGCGNGNMFASQKECLQTCRTVAA-- 60  
QY 69 TGLATSRNAADSVPSAPRRQDSHSDMFNFEYCTANAVTGPCRASFFRWYFDVER 128  
Db 61 -----CNLPVIRGPCRAFIQLWAFDAVK 83  
QY 129 NSCNFIYGGCGNKNYSRSEACMLRC 156  
Db 84 GKCVLPYGGCGNGNKNFYSEKCEYIC 111  
RESULT 12  
5223482-22  
; Patent No. 5223482  
; APPLICANT: SCHILLING, JAMES W.; PONTE, PHYLLIS A.; CORDELL,  
; BARBARA  
; TITLE OF INVENTION: RECOMBINANT ALZHEIMER'S PROTEASE  
; INHIBITORY AMYLOID PROTEIN AND METHOD OF USE  
; NUMBER OF SEQUENCES: 34  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/07/361,912  
; FILING DATE: 06-JUN-1989  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 359,911  
; FILING DATE: 12-MAY-1989  
; APPLICATION NUMBER: 87,002  
; FILING DATE: 18-AUG-1987  
; APPLICATION NUMBER: 8,810  
; FILING DATE: 30-JAN-1987  
; APPLICATION NUMBER: 948,376  
; FILING DATE: 31-DEC-1986  
; APPLICATION NUMBER: 932,193  
; FILING DATE: 17-NOV-1986  
; SEQ ID NO: 22:  
; LENGTH: 122  
5223482-22

Query Match 24.6%; Score 233.5; DB 5; Length 122;  
Best Local Similarity 33.6%; Pred. No. 1.8e-17;  
Matches 44; Conservative 15; Mismatches 33; Indels 39; Gaps 1;  
QY 26 RWTNTVDGSCQLFYVGGCGNSNNYLTKEBCLKKCATVTENATGLATSRNAADSVPS 85  
Db 22 RYFYNGTSMACETFYGGCGMGNLNFSLQKECLQTCRTV----- 60  
QY 86 APRQDSEHSDMFNFEYCTANAVTGPCRASFFRWYFDVERNSCNNFIYGGCGRNKNS 145  
Db 61 -----EACNLPVIRGPCRAFIQLWAFDAVKGVKCVRFSGYGGCKGNGK 102  
QY 146 YRSEACMLRC 156  
Db 103 FYSQKECEYIC 113

RESULT 13  
US-07-828-920A-1  
; Sequence 1, Application US/07828920A  
; Patent No. 5312736  
; GENERAL INFORMATION:  
; APPLICANT: Rasmussen, Jesper  
; TITLE OF INVENTION: Anticoagulant Protein  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: No. 5312736 No. 5312736disk of No. 5312736th America, Inc.  
; STREET: 405 Lexington Avenue, Suite 6200  
; CITY: New York  
; STATE: New York  
; COUNTRY: United States of America  
; ZIP: 10174-6201

```
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/828,920A
FILING DATE: 19920127
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DK 4080/89
FILING DATE: 18-AUG-1989
APPLICATION NUMBER: WO PCT/DK90/00212
FILING DATE: 17/AUG/1990
ATTORNEY/AGENT INFORMATION:
NAME: Zelson, Steve T.
REGISTRATION NUMBER: 30335
REFERENCE/DOCKET NUMBER: 3287.204-US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212 867 0123
TELEFAX: 212 867 0298
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: AMINO ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
ORIGINAL SOURCE:
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: Protein
LOCATION: 1..276
US-07-828-920A-1

Query Match 24.6%; Score 233.5; DB 1; Length 276;
Best Local Similarity 33.6%; Pred. No. 5.1e-17;
Matches 51; Conservative 26; Mismatches 64; Indels 11; Gaps 3;

Qy 9 DFCIVSKVGRCRASPRWYNNVTDGSQLFYGGCDGNSNNYLTKECKLKCAVTENA 68
||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| :
Db 95 DFCFLEDPGICRGYITRYFNNQTKQCFKYGCGLGNNMNFETLECKNIC---EDGP 151
| : | : | : | : | : | : | : | : | : | : | : | : | : | : | : | :
Qy 69 TG----DLATSRNAADSVPSAPRQDSHSDMFNFEYECTANAVTGPCRASFPRWYF 124
| : | : | : | : | : | : | : | : | : | : | : | : | : | : | : | :
Db 152 NGFQVDNYGTQLNAVNSLTP---QSTKVPSEFHFHGPSNCLTPADRGGLCRANENRFY 207
: : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
125 DVERNSCNFIYGGCGNGKNSYSEACMLRC 156
208 NSVIGKCRPFYSGCGGNNENFTSKQECILRAC 239

RESULT 14
US-08-437-841-9
Sequence 9, Application US/08437841
Patent No. 5563123
GENERAL INFORMATION:
APPLICANT: Innis, Michael
APPLICANT: Creasey, Abia
TITLE OF INVENTION: Chimeric Proteins
NUMBER OF SEQUENCES: 37
CORRESPONDENCE ADDRESS:
ADDRESSEE: Chiron Corporation
STREET: 4560 Horton St.
CITY: Emeryville
STATE: CA
COUNTRY: USA
ZIP: 94608
COMPUTER READABLE FORM:
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/828,920A
FILING DATE: 05-AUG-1994
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Savereide, Paul B.
REGISTRATION NUMBER: 36,914
REFERENCE/DOCKET NUMBER: 0990.001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 510-601-2585
```

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SOFTWARE: PatentIn Release #1.0, Version #1.30B
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/437,841
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/286,521
FILING DATE: 05-AUG-1994
ATTORNEY/AGENT INFORMATION:
NAME: Savereide, Paul B.
REGISTRATION NUMBER: 36,914
REFERENCE/DOCKET NUMBER: 0990.001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 510-601-2585
TELEFAX: 510-655-3542
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-437-841-9

Query Match 24.6%; Score 233.5; DB 1; Length 276;
Best Local Similarity 33.6%; Pred. No. 5.1e-17;
Matches 51; Conservative 26; Mismatches 64; Indels 11; Gaps 3;

Qy 9 DFCIVSKVGRCRASPRWYNNVTDGSQLFYGGCDGNSNNYLTKECKLKCAVTENA 68
||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| : ||| :
Db 95 DFCFLEDPGICRGYITRYFNNQTKQCFKYGCGLGNNMNFETLECKNIC---EDGP 151
| : | : | : | : | : | : | : | : | : | : | : | : | : | : | : | :
Qy 69 TG----DLATSRNAADSVPSAPRQDSHSDMFNFEYECTANAVTGPCRASFPRWYF 124
| : | : | : | : | : | : | : | : | : | : | : | : | : | : | : | :
Db 152 NGFQVDNYGTQLNAVNSLTP---QSTKVPSEFHFHGPSNCLTPADRGGLCRANENRFY 207
: : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
125 DVERNSCNFIYGGCGNGKNSYSEACMLRC 156
208 NSVIGKCRPFYSGCGGNNENFTSKQECILRAC 239

RESULT 15
US-08-286-521-9
Sequence 9, Application US/08286521
Patent No. 5589359
GENERAL INFORMATION:
APPLICANT: Innis, Michael
APPLICANT: Creasey, Abia
TITLE OF INVENTION: Chimeric Proteins
NUMBER OF SEQUENCES: 37
CORRESPONDENCE ADDRESS:
ADDRESSEE: Chiron Corporation
STREET: 4560 Horton St.
CITY: Emeryville
STATE: CA
COUNTRY: USA
ZIP: 94608
COMPUTER READABLE FORM:
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30B
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/286,521
FILING DATE: 05-AUG-1994
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Savereide, Paul B.
REGISTRATION NUMBER: 36,914
REFERENCE/DOCKET NUMBER: 0990.001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 510-601-2585
```

TELEFAX: 510-655-3542  
INFORMATION FOR SEQ ID NO: 9:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 276 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
US-08-286-521-9

Query Match 24.6%; Score 233.5; DB 1; Length 276;  
Best Local Similarity 33.6%; Pred. No. 5.1e-17;  
Matches 51; Conservative 26; Mismatches 64; Indels 11; Gaps 3;  
QY 9 DFCLVSKVGRCRASMPRWYNNVTDGSCQLFYGGCDGNSNNYLTKEECKLKCATVTENA 68  
| | | : : | | | : | : | | | | | | | | :  
Db 95 DFCFLEEDPGICRGYITRYFYNNQTKQCFERKYGGCLGNMNNFETLECKNIC---EDGP 151  
69 TG----DLATSRNADSSVPSAPRRQDSHSDMFNYEYCTANAVTGPCRASFPRWYF 124  
| : | | | : | : | | : | : | | | | : | :  
Db 152 NGFQVDNYGTQLNAVNNSLTP----QSTKVPSLFEFHGPSWCLTPADRGLCRANENRFY 207  
QY 125 DVERNSCNNFIYGGCRGKNKNSYRSEACMLRC 156  
| | | | | | | | | | | : | : | : | : | : |  
Db 208 NSVIGKCRPFKYGGCGGNNENFTSKQECRLAC 239

Search completed: January 31, 2001, 15:05:21  
Job time: 113 sec





GenCore version 4.5  
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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:05:08 ; Search time 15.54 Seconds  
(without alignments)  
742.800 Million cell updates/sec

Title: US-09-441-654A-1

Perfect score: 170

Sequence: 1 ADERSIHDFCLVSKVYGRG.....ACMLRCFRQENPPLPLGSK 170

Scoring table:

OLIGO  
Gapop 60.0 , Gapext 60.0  
195891 seqs, 67900655 residues

id size : 0

Total number of hits satisfying chosen parameters: 195891

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : PIR66:\*

1: pir1.\*

2: pir2.\*

3: pir3.\*

4: pir4.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	13	7.6	252	JG0185	hepatocyte growth
2	11	6.5	302	TIRTK	tissue factor path
3	9	5.3	58	TIRABK	isoactivator K (BP
4	9	5.3	65	TIVIVC	venom basic protei
5	8	4.7	922	T23573	hypothetical prote
6	8	4.7	973	S54534	coatmer complex b
7	8	4.7	1599	T16210	hypothetical prote
8	8	4.7	2844	S28291	hypothetical prote
9	7	4.1	57	TIFHBP	proteinase inhibit
10	7	4.1	61	TIRCBP	proteinase inhibit
11	7	4.1	62	S19327	venom basic protei
12	7	4.1	120	JQ1280	lipid transfer pro
13	7	4.1	125	TIRHBI	alpha-1-microglobu
14	7	4.1	183	T28711	hypothetical prote
15	7	4.1	210	S66484	insulin-like growt
16	7	4.1	253	S49183	hypothetical prote
17	7	4.1	438	T12494	hypothetical prote
18	7	4.1	500	F71978	hypothetical prote
19	7	4.1	507	H82378	probable long-chain
20	7	4.1	988	I50611	protein-tyrosine k
21	7	4.1	1663	C3HU	complement C3 prec
22	7	4.1	2167	T34395	hypothetical prote
23	7	4.1	2172	T20145	hypothetical prote
24	7	4.1	4660	T42737	gp330 protein prec
25	6	3.5	55	S30332	proteinase inhibit
26	6	3.5	56	JN0380	trypsin inhibitor
27	6	3.5	57	TIRIVC	venom basic protei
28	6	3.5	57	TINJVC	venom basic protei
29	6	3.5	57	S13846	venom animal kunit

#### ALIGNMENTS

##### RESULT 1

JG0185 hepatocyte growth factor activator inhibitor type 2 - mouse

C:Species: Mus musculus (House mouse)

C:Date: 23-Jul-1999 #sequence\_revision 23-Jul-1999 #text\_change 11-May-2000

C:Accession: JG0185

R:Ittoh, H.; Kataoka, H.; Hamasuna, R.; Kitamura, N.; Koono, M.

Biochem. Biophys. Res. Commun. 255, 740-748, 1999

A:Title: Hepatocyte growth factor activator inhibitor type 2 lacking the first

A:Reference number: JG0185; MUID:99160423

A:Accession: JG0185

A>Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-252 <IIO>

A:Cross-references: GB:AF099016

C:Superfamily: animal Kunitz-type proteinase inhibitor homology

F:133-183/Domain: animal Kunitz-type proteinase inhibitor homology <BPI>

Query Match 7.6% Score 13; DB 2; Length 252;

Best Local Similarity 100.0%; Pred. No. 5.3e-06;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 FIYGGCRGNKNSY 146

Db 161 FIYGGCRGNKNSY 173

|||||

|||||

##### RESULT 2

TIRTK tissue factor pathway inhibitor precursor - rat

N:Alternate names: extrinsic pathway inhibitor; lipoprotein-associated

C:Species: Rattus norvegicus (Norway rat)

C:Date: 30-Sep-1993 #sequence\_revision 30-Sep-1993 #text\_change 16-

C:Accession: JX0213

R:Enyioji, K.; Emi, M.; Mukai, T.; Kato, H.

J. Biochem. 111, 681-687, 1992

A:Title: cDNA cloning and expression of rat tissue factor pathway

A:Reference number: JX0213; MUID:92348361

A:Accession: JX0213

A:Molecule type: mRNA

A:Residues: 1-302 <ENJ>

A:Cross-references: DDBJ:D10926; NID:g220916; PIDN:BAA01724.1; PID:g220917

A:Experimental source: liver

C:Comment: This serine proteinase inhibitor regulates clotting by factor Xa-

C:Comment: The first Kunitz-type domain binds the factor VIIa/tissue factor comp-

C:Superfamily: tissue factor pathway inhibitor; animal Kunitz-type proteinase inh-

F:Keywords: anticoagulant; blood coagulation; duplication; glycoprotein; heparin

F:1-28/Domain: signal sequence #status predicted <Sig>

F:29-302/Product: tissue factor pathway inhibitor #status predicted <MAT>

F:53-103/Domain: animal Kunitz-type proteinase inhibitor homology <BPI>

F:124-174/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>

basic proteinase i  
isoaprotinin G2 -  
isoaprotinin G1 -  
serum basic protei  
hypothetical prote  
venom basic protei  
proteinase inhibit  
chymotrypsin inhib  
chymotrypsin inhib  
taicetoxin serine  
acrosin inhibitor  
chymotrypsin inhib  
9K protein - vacci  
Alzheimer's diseas  
Alzheimer's diseas  
Alzheimer's diseas

30 6 3.5 57 2 A59204  
31 6 3.5 58 2 S10063  
32 6 3.5 59 2 S00371  
33 6 3.5 60 1 TIBOR  
34 6 3.5 60 2 D81898  
35 6 3.5 61 1 TIVIT1  
36 6 3.5 62 2 S07451  
37 6 3.5 62 2 S01802  
38 6 3.5 62 2 S01803  
39 6 3.5 62 2 A44180  
40 6 3.5 63 1 TIFEF  
41 6 3.5 63 1 TIMCS  
42 6 3.5 76 1 WWVZRO  
43 6 3.5 76 2 S04855  
44 6 3.5 76 2 S03607  
45 6 3.5 76 2 S06678

F:222-272/Domain: animal Kunitz-type proteinase inhibitor homology <BP3>  
 F:288-291/Region: heparin binding #status predicted  
 F:53-103/62-86,78-99,124-174,133-157,149-170,222-272,231-255,247-268/Disulfide bonds: #s  
 F:63/inhibitory site: Lys (coagulation factor VII/tissue factor complex) #status predicted  
 F:134/inhibitory site: Arg (coagulation factor X) #status predicted  
 F:144,251,261/Binding site: carboxylate (Asn) (covalent) #status predicted  
 F:232/inhibitory site: Lys (unidentified proteinase) #status predicted

Query Match 6.5% Score 11; DB 1; Length 302;

Best Local Similarity 100.0%; Pred. No. 0.00084; Mismatches 0; Indels 0; Gaps 0;

Matches 11; Conservative 0;

QY 134 FIYGGCRGNK 144

Db 81 FIYGGCRGNK 91

RESULT 3

TIHABK

A:Title: Inhibitor K (BPI type) - Roman snail

A:Species: Helix pomatia (Roman snail)

C:Date: 23-Oct-1981 #sequence\_revision 23-Oct-1981 #text\_change 05-Aug-1994

C:Accession: A91232; A01225

R:Tschesche, H.; Dietl, T.

Eur J. Biochem. 58:439-451, 1975

A:Title: The amino-acid sequence of isoinhibitor K from snails (Helix pomatia). A sequen

A:Reference number: A91232; MUID:76043680

A:Accession: A91232

A:Molecule type: protein

A:Residues: 1-58 <TSC>

R:Dietl, T.; Tschesche, H.

Hoppe-Seyler's Z. Physiol. Chem. 357, 139-145, 1976

A:Title: Die Disulfidbruecken des Trypsin-Kallikrein-Inhibitors K aus Weinbergschnecken

A:Reference number: A91666; MUID:76141310

A:Contents: annotation: disulfide bonds

C:Comment: This is one of several isoinhibitors of broad specificity that are secreted i

C:Superfamily: basic proteinase inhibitor; animal Kunitz-type proteinase inhibitor homol

C:Keywords: pyroglutamic acid; serine proteinase inhibitor

F:7-57/Domain: animal Kunitz-type proteinase inhibitor homology <BPI>

F:1/Modified site: pyrrolidone carboxylic acid (Gln) #status experimental

F:7-57,16-40,32-53/Disulfide bonds: #status predicted

Query Match 5.3% Score 9; DB 1; Length 58;

Best Local Similarity 100.0%; Pred. No. 0.027; Mismatches 0; Indels 0; Gaps 0;

Matches 9; Conservative 0;

QY 134 FIYGGCRGN 142

Db 35 FIYGGCRGN 43

RESULT 4

TIVIVC

venom basic proteinase inhibitor III - sand viper

N:Alternate names: venom chymotrypsin inhibitor

C:Species: Vipera ammodytes (sand viper)

C:Date: 17-May-1985 #sequence\_revision 17-May-1985 #text\_change 16-Aug-1996

C:Accession: A01223

R:Bitonja, A.; Meloun, B.; Gubensek, F.

Biochim. Biophys. Acta 746, 138-145, 1983

A:Title: The primary structure of Vipera ammodytes venom chymotrypsin inhibitor.

A:Reference number: A01223

A:Accession: A01223

A:Molecule type: protein

A:Residues: 1-65 <RT>

C:Superfamily: basic proteinase inhibitor; animal Kunitz-type proteinase inhibitor homol

C:Keywords: serine proteinase inhibitor; venom

F:7-57/Domain: animal Kunitz-type proteinase inhibitor homology <BPI>

F:7-57,16-40,32-53/Disulfide bonds: #status predicted

F:17/inhibitory site: Leu (chymotrypsin) #status predicted

Query Match 5.3% Score 9; DB 1; Length 65;

Best Local Similarity 100.0%; Pred. No. 0.03; Mismatches 0; Indels 0; Gaps 0;

Matches 9; Conservative 0;

QY 134 FIYGGCRGN 142

Db 35 FIYGGCRGN 43

RESULT 5

T23573

hypothetical protein:K10D3.4 - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C:Date: 15-Oct-1999 #sequence\_revision 15-Oct-1999 #text\_change 15-Oct-1999

C:Accession: T23573

R:McMurray, A.

submitted to the EMBL Data Library, June 1996

A:Reference number: Z19762

A:Accession: T23573

A:Status: preliminary; translated from GB/EMBL/DDBJ

A:Molecule type: DNA

A:Residues: 1-922 <N1L>

A:Cross-references: EMBL:275545; PIDN:CAA99886.1; GSPDB:GNO0019; CESP:K10D3.4

A:Experimental source: clone K10D3

C:Genetics:

A:Gene: CESP:K10D3.4

A:Map position: 1

A:Introns: 60/1; 228/1; 278/1; 355/1; 743/1; 802/1; 885/2

Query Match 4.7% Score 8; DB 2; Length 922;

Best Local Similarity 100.0%; Pred. No. 3.4; Mismatches 0; Indels 0; Gaps 0;

Matches 8; Conservative 0;

QY 43 GCDGNSNN 50

Db 552 GCDGNSNN 559

RESULT 6

S54534

coatomer complex beta chain - yeast (Saccharomyces cerevisiae)

N:Alternate names: protein YB8419.05c; protein YDR238C

C:Species: Saccharomyces cerevisiae

C:Date: 08-Jul-1995 #sequence\_revision 01-Sep-1995 #text\_change 29-Sep-1999

C:Accession: S54534; A55123; C55123; S50260

R:Oliver, K.; Harris, D.

submitted to the EMBL Data Library, May 1995

A:Reference number: S54534

A:Accession: S54534

A:Molecule type: DNA

A:Residues: 1-973 <OLI>

A:Cross-references: EMBL:249701; NID:9817819; PIDN:CAA89724.1; PID:9817824; MIPS:1000

A:Experimental source: strain AB972

R:Duden, R.; Hosobuchi, M.; Hamamoto, S.; Winey, M.; Byers, B.; Schekman, R.

J. Biol. Chem. 269, 24486-24495, 1994

A:Title: Yeast beta'- and beta''-coat proteins (COP). Two coatomer subunits essential

A:Reference number: A55123; MUID:95014199

A:Accession: A55123

A:Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-411, E, 413-973 <UUD>

A:Cross-references: GB:U11236; NID:9595412; PIDN:AAA61710.1; PID:9595413

A:Accession: C55123

A:Molecule type: protein

A:Residues: 353-362,496-514;645-655;934-942 <DU2>

C:Genetics:

A:Gene: SGD:SGC26

A:Cross-references: SGD:S0002646; MIPS:YDR238C

A:Map position: 4R

C:Superfamily: coatomer complex beta chain

C:Keywords: blocked amino end; transmembrane protein

F:391-407/Domain: transmembrane #status predicted <TM1>  
F:409-425/Domain: transmembrane #status predicted <TM2>  
F:587-603/Domain: transmembrane #status predicted <TM3>

Query Match 4.7%; Score 8; DB 2; Length 973;  
Best Local Similarity 100.0%; Pred. No. 3.6;  
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 71 DLATSRNA 78  
Db 347 DLATSRNA 354  
|||||

RESULT 7  
Ti6210  
hypothetical protein F30H5.3 - Caenorhabditis elegans  
C:Species: Caenorhabditis elegans  
Date: 20-Sep-1999 #sequence\_revision 20-Sep-1999 #text\_change 20-Sep-1999  
Accession: T16210  
Pauley, A.; Stelliyes, L.  
submitted to the EMBL Data Library, June 1995

A:Description: The sequence of C. elegans cosmid F30H5.  
A:Reference number: Z18478  
A:Accession: T16210  
A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA  
A:Residues: 1-1599 <PAU>  
A:Cross-references: EMBL:U29096; NID:g861390; PID:g861393; PIDN:AAA68408.1; CESP:F30H5.3  
A:Experimental source: strain Bristol N2  
C:Genetics:  
A:Gene: CESP:F30H5.3  
A:Introns: 12/1; 59/2; 85/3; 124/3; 217/2; 534/3; 560/1; 1549/1

Query Match 4.7%; Score 8; DB 2; Length 1599;  
Best Local Similarity 100.0%; Pred. No. 5.6;  
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 GCDGNSNN 50  
Db 704 GCDGNSNN 711  
|||||

RESULT 8  
S28291  
hypothetical protein ZC84.1 - Caenorhabditis elegans  
C:Species: Caenorhabditis elegans  
Date: 13-Jan-1995 #sequence\_revision 13-Jan-1995 #text\_change 29-Aug-1997  
Accession: S28291  
R:Thomas, K.

submitted to the EMBL Data Library, December 1992  
A:Reference number: S28285  
A:Accession: S28291  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-2844 <THO>  
A:Cross-references: EMBL:Z19157  
C:Genetics:  
A:Introns: 14/1; 32/3; 57/1; 192/3; 277/1; 398/1; 439/1; 474/1; 497/1; 813/1; 1135/1; 1249/1; 2555/1; 2720/1; 2739/3; 2819/1

C:Superfamily: animal Kunitz-type proteinase inhibitor homology  
F:220-274/Domain: animal Kunitz-type proteinase inhibitor homology <BPI1>  
F:343-395/Domain: animal Kunitz-type proteinase inhibitor homology <BPI2>  
F:442-492/Domain: animal Kunitz-type proteinase inhibitor homology <BPI3>  
F:546-598/Domain: animal Kunitz-type proteinase inhibitor homology <BPI4>  
F:654-706/Domain: animal Kunitz-type proteinase inhibitor homology <BPI5>  
F:1662-1716/Domain: animal Kunitz-type proteinase inhibitor homology <BPI6>  
F:1787-1839/Domain: animal Kunitz-type proteinase inhibitor homology <BPI7>  
F:1845-1895/Domain: animal Kunitz-type proteinase inhibitor homology <BPI8>  
F:1952-2004/Domain: animal Kunitz-type proteinase inhibitor homology <BPI9>  
F:2097-2152/Domain: animal Kunitz-type proteinase inhibitor homology <BPI10>

Query Match 4.7%; Score 8; DB 2; Length 2844;  
Best Local Similarity 100.0%; Pred. No. 9.1;  
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 GCDGNSNN 50  
Db 580 GCDGNSNN 587  
|||||

RESULT 9  
TIFHBP  
proteinase inhibitor - flesh fly (Sarcophaga bullata)  
C:Species: Sarcophaga bullata  
C:Date: 07-Feb-1992 #sequence\_revision 22-Jul-1994 #text\_change 07-May-1993  
C:Accession: A37294  
R:Papayannopoulos, I.A.; Blemann, K.  
Protein Sci. 1, 278-288, 1992  
A:Title: Amino acid sequence of a protease inhibitor isolated from Sarcophaga  
A:Reference number: A37294; MUID:93284121  
A:Accession: A37294  
A:Molecule type: protein  
A:Residues: 1-57 <PAP>  
A:Experimental source: hemolymph  
C:Superfamily: basic proteinase inhibitor; animal Kunitz-type proteinase  
C:Keywords: serine proteinase inhibitor  
F:6-56/Domain: animal Kunitz-type proteinase inhibitor homology <BPI>  
F:6-56,15-39,31-52/Disulfide bonds: #status predicted  
F:16/inhibitory site: Arg (chymotrypsin) #status predicted

Query Match 4.1%; Score 7; DB 1; Length 57;  
Best Local Similarity 100.0%; Pred. No. 3.6;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 136 YGGCRGN 142  
Db 36 YGGCRGN 42  
|||||

RESULT 10  
TIHCBP  
proteinase inhibitor (BPI-type) - horseshoe crab (Tachyplesus tridentatus)  
C:Species: Tachyplesus tridentatus  
C:Date: 08-Mar-1989 #sequence\_revision 22-Jul-1994 #text\_change 24-Feb-1995  
C:Accession: A26923  
R:Nakamura, T.; Hirai, T.; Tokunaga, F.; Kawabata, S.; Iwanaga, S.  
J. Biochem. 101, 1297-1306, 1987  
A:Title: Purification and amino acid sequence of Kunitz-type protease inhibitor found  
A:Reference number: A26923; MUID:88007472  
A:Accession: A26923  
A:Molecule type: protein  
A:Residues: 1-61 <NAK>  
A:Experimental source: hemocytes  
C:Comment: The inhibitory activity is similar to bovine basic proteinase inhibitor.  
C:Superfamily: basic proteinase inhibitor; animal Kunitz-type proteinase inhibitor  
C:Keywords: serine proteinase inhibitor  
F:8-58/Domain: animal Kunitz-type proteinase inhibitor homology <BPI>  
F:8-58,17-41,33-54/Disulfide bonds: #status predicted  
F:18/inhibitory site: Arg (chymotrypsin, elastase, trypsin, plasmin, kallikrein)

Query Match 4.1%; Score 7; DB 1; Length 61;  
Best Local Similarity 100.0%; Pred. No. 3.8;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 111 VTGPCRA 117  
Db 13 VTGPCRA 19  
|||||

RESULT 11  
S19327

venom basic proteinase inhibitor - leaf-nosed viper  
N:Alternate names: trypsin inhibitor (Kunitz-type)  
C:Species: Eristocophis machahoni (leaf-nosed viper)  
C:Date: 22-Nov-1993 #sequence\_revision 03-Nov-1995 #text\_change 16-Jul-1999  
A:Accession: S19327  
R:Siddiqui, A.R.; Zaidi, Z.H.; Joernvall, H.  
FEBS Lett. 294, 141-143, 1991  
A:Title: Purification and characterization of a Kunitz-type trypsin inhibitor from Leaf-  
A:Reference number: S19327; MUID:92077130  
A:Accession: S19327  
A:Status: preliminary  
A:Molecule type: mRNA  
A:Residues: 1-62 <SID>  
C:Superfamily: basic proteinase inhibitor; animal Kunitz-type proteinase inhibitor homol  
C:Keywords: serine proteinase inhibitor; venom  
F:2-52/Domain: animal Kunitz-type proteinase inhibitor homology <BP1>  
  
Query Match 4.1%; Score 7; DB 2; Length 62;  
Best Local Similarity 100.0%; Pred. No. 3.9;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 133 NFIYGGC 139  
DB 29 NFIYGGC 35  
|||||||  
RESULT 12  
JQ1280  
lipid transfer protein EP2 precursor - carrot  
N:Alternate names: extracellular protein 2  
C:Species: Daucus carota (carrot)  
C:Date: 10-Sep-1999 #sequence\_revision 10-Sep-1999 #text\_change 10-Sep-1999  
A:Accession: JQ1280  
R:Sterk, P.; Boelj, H.; Schellekens, G.A.; Van Kammen, A.; De Vries, S.C.  
Plant Cell 3, 907-921, 1991  
A:Title: Cell-specific expression of the carrot EP2 lipid transfer protein gene.  
A:Reference number: JQ1280; MUID:92361243  
A:Accession: JQ1280  
A:Molecule type: mRNA  
A:Residues: 1-120 <STE>  
A:Cross-references: GB:M64746; NID:g167553; PIDN:AAB96834.1; PID:g167554  
C:Comment: This protein locates in cell walls.  
C:Comment: The gene encoding for this protein is expressed in the embryogenic cell cultu  
C:Superfamily: phospholipid transfer protein  
F:1-25/Domain: signal sequence #status predicted <SIG>  
F:27-120/Product: lipid transfer protein EP2 #status predicted <LIP>  
  
Query Match 4.1%; Score 7; DB 1; Length 120;  
Best Local Similarity 100.0%; Pred. No. 6.9;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 107 TANAVTG 113  
DB 83 TANAVTG 89  
|||||||  
RESULT 13  
TIHOBI  
alpha-1-microglobulin/inter-alpha-trypsin inhibitor - horse (fragment)  
N:Alternate names: EI-14 (inhibitory fragment of ITI); ITI; trypsin inhibitor, E-UTI  
C:Species: Equus caballus (domestic horse)  
C:Date: 30-Jun-1987 #sequence\_revision 04-Feb-2000 #text\_change 05-May-2000  
A:Accession: A01210; A45653  
R:Hochstrasser, K.; Wächter, E.; Albrecht, G.J.; Reisinger, P.  
Biol. Chem. Hoppe-Seyler 366, 473-478, 1985  
A:Title: Kunitz-type proteinase inhibitors derived by limited proteolysis of the inter-a  
A:Reference number: A06885; MUID:85225967  
A:Accession: A01210  
A:Molecule type: protein  
A:Residues: 3-125 <HOC>  
R;Veeraragavan, K.; Singh, K.; Wächter, E.; Hochstrasser, K.

Biochem. Int. 26, 405-413, 1992  
A:Title: Characterization of a trypsin inhibitor from equine urine.  
A:Reference number: A45653; MUID:92328813  
A:Accession: A45653  
A:Status: preliminary  
A:Molecule type: protein  
A:Residues: 1-12, 'E', 14-33 <VEE>  
A:Cross-references: PIDN:AAB22430.1; PID:g250858  
A:Experimental source: urine  
A:Note: sequence extracted from NCBI backbone (NCBIP:107966)  
C:Comment: This inhibitory fragment, released from native ITI after limited proteolysis  
C:Comment: first domain interacts weakly with PMN-granulocytic elastase and not at all with panc  
C:Comment: The amino acid at position P2' (19-Met) appears to determine the specificity  
C:elastase; those with leucine interact strongly.  
C:Superfamily: protein HC; animal Kunitz-type proteinase inhibitor homology  
C:Keywords: duplication; glycoprotein; plasma; serine proteinase inhibitor  
F:7-57/Domain: animal Kunitz-type proteinase inhibitor homology <BP2>  
F:63-113/Domain: animal Kunitz-type proteinase inhibitor homology  
F:7-57,16-40,32-53,63-113,72-96,88-109/disulfide bonds: #status predicted  
F:17/Inhibitory site: Leu (chymotrypsin, elastase) #status predicted  
F:26/Binding site: carbohydrate (Asn) (covalent) #status experimental  
F:73/Inhibitory site: Arg (trypsin) #status predicted

Query Match 4.1%; Score 7; DB 1; Length 125;  
Best Local Similarity 100.0%; Pred. No. 7.1;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 136 YGGCRGN 142

DB 93 YGGCRGN 99

RESULT 14

T28711

hypothetical protein T21D12.12 - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C:Date: 15-Oct-1999 #sequence\_revision 15-Oct-1999 #text\_change 29-Oct-1999

C:Accession: T28711

R:Woessner, J.

submitted to the EMBL Data Library, August 1997

A:Description: The sequence of C. elegans cosmid T21D12.

A:Reference number: Z20514

A:Accession: T28711

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-183 <WOE>

A:Cross-references: EMBL:AF016687; PIDN:AAC48097.1; GSPDB:GN00022; CESP:T21D12.12

A:Experimental source: strain Bristol N2; clone T21D12

C:Genetics:

A:Gene: CESP:T21D12.12

A:Map position: 4

A:Introns: 57/3; 88/1; 126/1; 147/2

Query Match 4.1%; Score 7; DB 2; Length 183;

Best Local Similarity 100.0%; Pred. No. 9.9;

Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 GCDGNSN 49

DB 58 GCDGNSN 64

RESULT 15

S66484

insulin-like growth factor II precursor - spiny dogfish

C:Species: Squalus acanthias (spiny dogfish)

C:Date: 28-Oct-1996 #sequence\_revision 13-Mar-1997 #text\_change 16-Jul-1999

C:Accession: S66484; S58053

R:Duguay, S.J.; Chan, S.J.; Mommsen, T.P.; Steiner, D.F.

FEBS Lett. 371, 69-72, 1995

A:Title: Divergence of insulin-like growth factors I and II in the elasmobranch.

A:Reference number: S66484; MUID:95394151  
A:Accession: S66484  
A:Molecule type: mRNA  
A:Residues: 1-210 <DUG>  
A:Cross-references: EMBL:250082; NID:g902732; PIDN:CRA90413.1; PID:g902733  
A:Experimental source: liver  
C:Superfamily: insulin  
C:Keywords: growth factor  
F:1-49/Domain: signal sequence #status predicted <SIG>  
F:50-210/Product: insulin-like growth factor II #status predicted <MAT>  
F:50-82/Domain: insulin chain B-like #status predicted <DOB>  
F:83-90/Domain: insulin connecting peptide C-like #status predicted <CPE>  
F:91-111/Domain: insulin chain A-like #status predicted <DOA>  
F:112-117/Domain: Peptide D #status predicted <DOD>  
F:118-210/Domain: carboxyl-terminal propeptide (E peptide) #status predicted <CHE>  
F:58-97,70-110,96-101/Disulfide bonds: #status predicted

Query Match 4.1%; Score 7; DB 2; Length 210;  
Best Local Similarity 100.0%; Pred. No. 11;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 VSKVGR 19

|||||

Db 78 VSKVGR 84

Search completed: January 31, 2001, 15:07:08  
Job time: 120 sec



GenCore version 4.5  
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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:05:52 ; Search time 10.05 Seconds  
(without alignments)  
546.268 Million cell updates/sec

Title: US-09-441-654A-1

Perfect score: 170

Sequence: 1 ADERSIHDFCLSVKVGRC.....ACMLRCFRQENPLPLGSK 170

Scoring table:  
OLIGO  
Gapop 60.0 , Gapext 60.0

Searched: 88757 seqs, 32294092 residues

Word size : 0

Total number of hits satisfying chosen parameters: 88757

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : SwissProt\_39.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	11	6.5	302	1	TFPI_RAT
2	9	5.3	58	1	ISIK_HELPO
3	9	5.3	65	1	IVB3_VIPAA
4	8	4.7	973	1	COPB_YEAST
5	8	4.7	1416	1	YN81_CABEL
6	7	4.1	57	1	SBPI_SARBU
7	7	4.1	61	1	IBPI_TACTR
8	7	4.1	62	1	IVBT_ERIMA
9	7	4.1	120	1	NLTP_DAUCA
10	7	4.1	123	1	IATR_HORSE
11	7	4.1	521	1	TRPE_BUCAL
12	7	4.1	1663	1	CO3_HUMAN
13	7	4.1	4660	1	LRP2_RAT
14	6	3.5	52	1	ISP2_GALME
15	6	3.5	55	1	ISH1_STOHE
16	6	3.5	55	1	ISH2_STOHE
17	6	3.5	56	1	ITR4_RADMA
18	6	3.5	57	1	IVB2_HENHA
19	6	3.5	57	1	IVB2_NAJNI
20	6	3.5	57	1	IVBT_NAJNA
21	6	3.5	60	1	IBPS_BOVIN
22	6	3.5	61	1	ITRS_RAT
23	6	3.5	61	1	IVB1_VIPAA
24	6	3.5	62	1	IP52_ANESU
25	6	3.5	62	1	ISCL_BOMMO
26	6	3.5	62	1	ISCL_BOMMO
27	6	3.5	63	1	IC53_BOMMO
28	6	3.5	63	1	IMAP_DROFU
29	6	3.5	69	1	CRPT_BOOMI
30	6	3.5	76	1	A4_MACMU
31	6	3.5	76	1	Y8K9_VACCV
32	6	3.5	87	1	A4_MACFA
33	6	3.5	94	1	S110_RAT

34 34 6 3.5 100 1 BPT1\_BOVIN  
35 35 6 3.5 100 1 BPT2\_BOVIN  
36 36 6 3.5 110 1 IBP\_CARCR  
37 37 6 3.5 115 1 YDG5\_SCHPO  
38 38 6 3.5 122 1 UPTI\_PIG  
39 39 6 3.5 123 1 IATR\_SHEEP  
40 40 6 3.5 145 1 RL13\_STACA  
41 41 6 3.5 150 1 RL15\_MYCGE  
42 42 6 3.5 160 1 Y964\_MYCTU  
43 43 6 3.5 162 1 YLR2\_EBV  
44 44 6 3.5 176 1 RL1X\_HUMAN  
45 45 6 3.5 176 1 RL1X\_RAT

#### ALIGNMENTS

RESULT 1  
TFPI\_RAT  
ID TFPI\_RAT STANDARD; PRT; 302 AA.  
AC Q02445;  
DT 01-JUL-1993 (Rel. 26, Created)  
DT 01-JUL-1993 (Rel. 26, Last sequence update)  
DT 30-MAY-2000 (Rel. 39, Last annotation update)  
DE TISSUE FACTOR PATHWAY INHIBITOR PRECURSOR (TFPI) (LIPOPROTEIN-  
DE ASSOCIATED COAGULATION INHIBITOR) (LACI) (EXTRINSIC PATHWAY INHIBITOR).  
DE (EPI).  
GN TFPI.  
OS Rattus norvegicus (Rat).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN-SPRAGUE-DAWLEY; TISSUE=LIVER;  
RX MEDLINE=92348361; PubMed=1639767;  
RA Enjoji K.-I., Emi M., Mukai T., Kato H.;  
RT "cDNA cloning and expression of rat tissue factor pathway inhibitor  
RT (TFPI).";  
RL J. Biochem. 111:681-687(1992).  
CC -!- FUNCTION: INHIBITS FACTOR X (X(A)) DIRECTLY AND, IN A XA-DEPENDENT  
WAY, INHIBITS VII(A)/TISSUE FACTOR ACTIVITY, PRESUMABLY BY FORMING  
A QUATERNARY X(A)/LACI/VII(A)/TF COMPLEX. IT POSSESSES AN  
ANTITHROMBOTIC ACTION AND ALSO THE ABILITY TO ASSOCIATE WITH  
LIPOPROTEINS IN PLASMA.  
CC -!- TISSUE SPECIFICITY: MOST ABUNDANT IN HEART, LUNG, KIDNEY, AND  
AORTIC ENDOTHELIAL CELLS.  
CC -!- DOMAIN: THIS INHIBITOR CONTAINS THREE INHIBITORY DOMAINS.  
CC -!- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
CC HIGHLY SIMILAR TO TFP2.  
-----  
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-----  
EMBL: D10926; BAA01724.1;  
PIR: JX0213; TIRTK.  
HSP: P10646; ITFX.  
DR INTERPRO: IPR002223;  
DR PFAM: PF00014; Kunitz\_BPTI; 3.  
DR PRINTS: PR00759; BASICPTASE.  
DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 3.  
DR PROSITE: PS50279; BPTI\_KUNITZ\_2; 3.  
KW Serine protease inhibitor; Glycoprotein; Repeat; Blood coagulation;  
Signal.  
FT CHAIN 1 28  
FT DOMAIN 29 302  
FT CHAIN 53 103  
FT DOMAIN 124 174  
FT TISSUE FACTOR PATHWAY INHIBITOR.  
FT BPTI/KUNITZ INHIBITOR 1  
FT (VII(A)/TISSUE FACTOR BINDING SITE).  
FT BPTI/KUNITZ INHIBITOR 2

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FT DOMAIN 222 272  
 FT DISULFID 53 103  
 FT DISULFID 62 86  
 FT DISULFID 78 99  
 FT DISULFID 64 64  
 FT ACT\_SITE 63 174  
 FT DISULFID 124 157  
 FT DISULFID 133 170  
 FT DISULFID 149 170  
 FT ACT\_SITE 134 135  
 FT DISULFID 222 272  
 FT DISULFID 231 255  
 FT DISULFID 247 268  
 FT ACT\_SITE 232 233  
 FT CARBOHYD 144 144  
 FT CARBOHYD 251 251  
 FT CARBOHYD 261 261  
 FT SEQUENCE 302 AA; 34554 MW; F9AE82130A24A59F CRC64;

Query Match 6.5%; Score 11; DB 1; Length 302;  
 Best Local Similarity 100.0%; Pred. No. 0.00035;  
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 134 FIYGGCRGN 144  
 Db 81 FIYGGCRGN 91

RESULT 2  
 ISIK\_HELP0  
 ID ISIK\_HELP0 STANDARD; PRT; 58 AA.  
 AC P00994;  
 DT 21-JUL-1986 (Rel. 01, Created)  
 DT 21-JUL-1986 (Rel. 01, Last sequence update)  
 DT 01-AUG-1990 (Rel. 15, Last annotation update)  
 DE ISOINHIBITOR K.  
 OS Helix pomatia (Roman snail) (Edible snail).  
 OC Eukaryota; Metazoa; Mollusca; Gastropoda; Stylommatophora;  
 OC Helicoidae; Helicidae; Helix.  
 RN [1]  
 RP SEQUENCE.  
 RX MEDLINE=76043680; PubMed=1183446;  
 RA Tschesche H., Dietl T.;  
 RT "The amino-acid sequence of isoinhibitor K form snails (Helix pomatia). A sequence determination by automated Edman degradation and mass-spectral identification of the phenylthiohydantoins.";  
 RT Eur. J. Biochem. 58:439-451(1975).  
 RL [2]

DISULFIDE BONDS.  
 MEDLINE=76141310; PubMed=3462;  
 Dietl T., Tschesche H.;  
 RT "The disulfide bridges of the trypsin-kallikrein inhibitor K from snails (Helix pomatia). Thermal inactivation and proteolysis by thermolysin.";  
 RT Hoppe-Sevler's Z. Physiol. Chem. 357:139-145(1976).  
 CC -1- FUNCTION: THIS IS ONE OF SEVERAL ISOINHIBITORS OF BROAD SPECIFICITY THAT ARE SECRETED INTO THE MUCUS OF THE SNAIL.  
 CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.

DR PIR; A01223; TIRASB.  
 DR HSSP; P00974; IRRB.  
 DR INTERPRO; IPR002223; -.  
 DR PFAM; PF00014; Kunitz\_BPTI; 1.  
 DR PRINTS; PR00759; BASICTPASE.  
 DR PROSITE; PS00280; BPTI\_KUNITZ\_1; 1.  
 DR PROSITE; PS00279; BPTI\_KUNITZ\_2; 1.  
 KW Serine protease inhibitor.  
 FT MOD\_RES 1 1 PYRROLIDONE CARBOXYLIC ACID.  
 FT DISULFID 7 57  
 FT DISULFID 16 40  
 FT DISULFID 32 53  
 FT ACT\_SITE 17 18  
 FT SEQUENCE 58 AA; 6451 MW; 6796586C488453B7 CRC64;

Query Match 5.3%; Score 9; DB 1; Length 58;  
 Best Local Similarity 100.0%; Pred. No. 0.011;  
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 FIYGGCRGN 142  
 Db 35 FIYGGCRGN 43

RESULT 3  
 IVB3\_VIPAA  
 ID IVB3\_VIPAA STANDARD; PRT; 65 AA.  
 AC P00992;  
 DT 21-JUL-1986 (Rel. 01, Created)  
 DT 21-JUL-1986 (Rel. 01, Last sequence update)  
 DT 01-MAY-1992 (Rel. 22, Last annotation update)  
 DE VENOM BASIC PROTEASE INHIBITOR III (VENOM CHYMOTRYPSIN INHIBITOR).  
 OS Vipera ammodytes ammodytes (Western sand viper).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Lepidosauria; Squamata; Scleroglossa; Serpentes; Colubroidea;  
 OC Viperidae; Viperinae; Vipera.  
 RN [1]  
 RP SEQUENCE.  
 RC TISSUE=VENOM;  
 RA Ritonja A., Meloun B., Gubensek F.;  
 RT "The primary structure of Vipera ammodytes venom chymotrypsin inhibitor.";  
 RL Biochim. Biophys. Acta 746:138-145(1983).  
 CC -1- FUNCTION: THIS PROTEIN INHIBITS CHYMOTRYPSIN.  
 CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 DR PIR; A01223; TIVIVC.  
 DR HSSP; P31713; ISHP.  
 DR INTERPRO; IPR002223; -.  
 DR PFAM; PF00014; Kunitz\_BPTI; 1.  
 DR PRINTS; PR00759; BASICTPASE.  
 DR PROSITE; PS00280; BPTI\_KUNITZ\_1; 1.  
 DR PROSITE; PS00279; BPTI\_KUNITZ\_2; 1.  
 KW Venom; Serine protease inhibitor.  
 FT DISULFID 7 57  
 FT DISULFID 16 40  
 FT DISULFID 32 53  
 FT ACT\_SITE 17 18  
 FT SEQUENCE 65 AA; 7556 MW; 9D526F8E3BF7CC57 CRC64;

Query Match 5.3%; Score 9; DB 1; Length 65;  
 Best Local Similarity 100.0%; Pred. No. 0.012;  
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 FIYGGCRGN 142  
 Db 35 FIYGGCRGN 43

RESULT 4  
 COPE\_YEAST  
 ID COPE\_YEAST STANDARD; PRT; 973 AA.  
 AC P41810; Q03779;  
 DT 01-NOV-1995 (Rel. 32, Created)  
 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 DT 01-NOV-1997 (Rel. 35, Last annotation update)  
 DE COATOMER BETA SUBUNIT (BETA-COAT PROTEIN) (BETA-COP).  
 GN SEC26 OR YDR238C OR YD8419.05C.  
 OS Saccharomyces cerevisiae (Baker's yeast).  
 OC Eukaryota; Fungi; Ascomycota; Saccharomycetes; Saccharomycetales;  
 OC Saccharomycetaceae; Saccharomyces.  
 RN [1]  
 RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.  
 RC STRAIN=RSY255;  
 RX MEDLINE=95014199; PubMed=7929113;  
 RA Duden R., Hosobuchi M., Hamamoto S., Winsey M., Byers B., Schekman R.;



RT "Yeast beta- and beta'-coat proteins (COP). Two coatomer subunits  
 RL essential for endoplasmic reticulum-to-Golgi protein traffic.";  
 RN J. Biol. Chem. 269:24486-24495(1994).  
 RP [2]  
 RC SEQUENCE FROM N.A.  
 RA STRAIN=S288C / AB972;  
 RL Oliver K., Harris D., Barrell B.G., Rajandream M.A.;  
 RL Submitted (MAY-1995) to the EMBL/GenBank/DBJ databases.  
 CC -!- FUNCTION: THE COATOMER IS A CYTOSOLIC PROTEIN COMPLEX THAT BINDS  
 CC TO DILYSINE MOTIFS AND REVERSIBLY ASSOCIATES WITH GOLGI NON-  
 CC CLATHRIN-COATED VESICLES, WHICH FURTHER MEDIANE BIOSYNTHETIC  
 CC PROTEIN TRANSPORT FROM THE ER, VIA THE GOLGI UP TO THE TRANS GOLGI  
 CC NETWORK. COATOMER COMPLEX IS REQUIRED FOR BUDDING FROM GOLGI  
 CC MEMBRANES, AND IS ESSENTIAL FOR THE RETROGRADE GOLGI-TO-ER  
 CC TRANSPORT OF DILYSINE-TAGGED PROTEINS (BY SIMILARITY).  
 CC -!- SUBUNIT: OLIGOMERIC COMPLEX THAT CONSISTS OF AT LEAST THE ALPHA,  
 CC BETA, BETA', GAMMA, DELTA, EPSILON AND ZETA SUBUNITS.  
 CC -!- SUBCELLULAR LOCATION: THE COATOMER IS CYTOPLASMIC OR POLYMERIZED  
 CC ON THE CYTOPLASMIC SIDE OF THE GOLGI, AS WELL AS ON THE  
 CC VESICLES/BUDS ORIGINATING FROM IT (BY SIMILARITY).  
 CC -!- PTM: THE N-TERMINUS IS BLOCKED.  
 CC -!- MISCELLANEOUS: BREFELDIN A INDUCES DISSOCIATION FROM THE GOLGI OF  
 CC THE BETA-COP AND PRESUMABLY THE OTHER COATOMER SUBUNITS (BY  
 CC SIMILARITY).  
 CC -!- SIMILARITY: SIGNIFICANT, OF THE N-TERMINAL HALF OF BETA-COP WITH  
 CC THOSE OF BETA-ADAPTINS.  
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 CC -----  
 CC EMBL; U11236; AAA61710.1; -;  
 DR EMBL; Z49701; CAA89724.1; -;  
 DR SGD; S0002646; SEC26.  
 \*KW Transport; Protein transport; Golgi stack; Membrane; Phosphorylation.  
 FT CONFLICT 412 412 D -> E (IN REF. 1).  
 SQ SEQUENCE 973 AA; 109019 MW; 885420DB026BCFA3 CRC64;  
 -----  
 Query Match 4.7%; Score 8; DB 1; Length 973;  
 Best Local Similarity 100.0%; Pred. No. 1.6;  
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 -----  
 Y 71 DLATSRNA 78  
 Db 347 DLATSRNA 354  
 |||||  
 YN81\_CAEEL STANDARD; PRT; 1416 AA.  
 AC Q03610;  
 DT 01-FEB-1994 (Rel. 28, Created)  
 DT 15-JUL-1998 (Rel. 36, Last sequence update)  
 DT 01-OCT-2000 (Rel. 40, Last annotation update)  
 DE HYPOTHETICAL 316.1 KDA PROTEIN ZC84.1 IN CHROMOSOME III.  
 GN ZC84.1.  
 OS Caenorhabditis elegans.  
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;  
 OC Rhabditidae; Peloderinae; Caenorhabditis.  
 [1]  
 SEQUENCE FROM N.A.  
 RC STRAIN=BRISTOL N2;  
 RX MEDLINE=94150718; PubMed=7906398;  
 RA Wilson R., Ainscough R., Anderson K., Baynes C., Berks M., Coulson A.,  
 RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Fraser A.,  
 RA Craxton M., Dear S., Du Z., Durbin R., Favello A., Fraser A.,  
 RA Fulton L., Gardner A., Green P., Hawkins T., Hillier L., Jier M.,  
 RA Johnston L., Jones M., Kershaw J., Kirsten J., Laister N.,

RA Latreille P., Lightning J., Lloyd C., Mortimore B., O'Callaghan M.,  
 RA Parsons J., Percy C., Rifken L., Roopra A., Saunders D., Showkeen R.,  
 RA Sims M., Smaldon N., Smith A., Smith M., Sonnhammer E., Staden R.,  
 RA Sulston J., Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K.,  
 RA Waterson R., Watson A., Weinstock L., Wilkinson-Sproat J.,  
 RA Wohldman P.,  
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of  
 RT elegans.";  
 RN Nature 368:32-38(1994).  
 RL [2]  
 RP REVISIONS.  
 RC STRAIN=BRISTOL N2;  
 RA Jones S.J.M.;  
 RL Submitted (NOV-1997) to the EMBL/GenBank/DBJ databases.  
 CC -!- SIMILARITY: STRONG, TO D1044.3.  
 CC -!- SIMILARITY: CONTAINS 5 PROTEASE INHIBITOR DOMAINS BELONGING TO THE  
 CC BPTI/KUNITZ FAMILY OF INHIBITORS.  
 CC -----  
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 CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).  
 CC -----  
 CC EMBL; Z19157; CAA79569.1; -;  
 DR PIR; S28291; S28291.  
 DR HSSP; P07204; ZADX.  
 DR WORMPEP; ZC84.1; CBI5020.  
 DR INTERPRO; IPR000561; -;  
 DR INTERPRO; IPR002223; -;  
 DR INTERPRO; IPR002899; -;  
 DR PFAM; PF01683; EB; 3.  
 DR PFAM; PF00014; Kunitz\_BPTI; 5.  
 DR PRINTS; PR00759; BASICPTASE.  
 DR PROSITE; PS00280; BPTI\_KUNITZ\_1; 3.  
 DR PROSITE; PS0279; BPTI\_KUNITZ\_2; 5.  
 DR PROSITE; PS01186; EGF\_2; UNKNOWN.1.  
 KW Hypothetical protein; Serine protease inhibitor; Repeat.  
 FT DOMAIN 212 266 BPTI/KUNITZ INHIBITOR.  
 FT DOMAIN 352 387 BPTI/KUNITZ INHIBITOR.  
 FT DOMAIN 434 484 BPTI/KUNITZ INHIBITOR.  
 FT DOMAIN 538 590 BPTI/KUNITZ INHIBITOR.  
 FT DOMAIN 646 698 BPTI/KUNITZ INHIBITOR.  
 SQ SEQUENCE 1416 AA; 152986 MW; 531CACELCB22F70D CRC64;  
 -----  
 Query Match 4.7%; Score 8; DB 1; Length 1416;  
 Best Local Similarity 100.0%; Pred. No. 2.3;  
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 -----  
 QY 43 GCDGNSNN 50  
 Db 572 GCDGNSNN 579  
 |||||  
 SBPI\_SARBU STANDARD; PRT; 57 AA.  
 AC P26228;  
 DT 01-NOV-1991 (Rel. 20, Created)  
 DT 01-NOV-1991 (Rel. 20, Last sequence update)  
 DT 01-NOV-1995 (Rel. 32, Last annotation update)  
 DE PROTEASE INHIBITOR (SBPI).  
 OS Sarcophaga bullata (Grey flesh fly) (Neobellieria bullata).  
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;  
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
 OC Cecidoidea; Sarcophagidae; Sarcophaga.  
 RN [1]  
 RP SEQUENCE.  
 RC TISSUE=LARVAL HEMOLYMPH;  
 RX MEDLINE=93284121; PubMed=1304909;

RA Papayannopoulos I.A., Biemann K.;  
 RT "Amino acid sequence of a protease inhibitor isolated from Sarcophaga  
 RL bullata determined by mass spectrometry.";  
 RL Protein Sci. 1:278-288(1992).  
 CC -!- FUNCTION: SEEMS TO INHIBITS TRYPSIN.  
 CC -!- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 DR PIR: A37294; A37294.  
 DR HSSP: P10646; 1ADZ.  
 DR INTERPRO: IPR002223; -.  
 DR PFAM: PF00014; Kunitz\_BPTI; 1.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 1.  
 DR PROSITE: PS0279; BPTI\_KUNITZ\_2; 1.  
 KW Serine protease inhibitor.  
 FT DISULFID 6 56 BY SIMILARITY.  
 FT DISULFID 15 39 BY SIMILARITY.  
 FT DISULFID 31 52 BY SIMILARITY.  
 FT ACT\_SITE 16 17 REACTIVE BOND (TRYPSIN) (BY SIMILARITY).  
 SQ SEQUENCE 57 AA; 6518 MW; FC512C5399E87241 CRC64;

Query Match 4.1%; Score 7; DB 1; Length 57;  
 Best Local Similarity 100.0%; Pred. No. 1.4;  
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 136 YGCGRGN 142

|||||

DB 36 YGCGRGN 42

## RESULT 7

IBPTI\_TACTR  
 ID IBPTI\_TACTR STANDARD; PRT; 61 AA.  
 AC P16044;  
 DT 01-APR-1990 (Rel. 14, Created)  
 DT 01-APR-1990 (Rel. 14, Last sequence update)  
 DT 01-AUG-1990 (Rel. 15, Last annotation update)  
 DE PROTEINASE INHIBITOR (BPTI-TYPE).  
 OS Tachypleus tridentatus (Japanese horseshoe crab).  
 OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Merostomata; Xiphosura;  
 OC Limulidae; Tachypleus.  
 RN [1]  
 RP SEQUENCE.

RC TISSUE=HEMOCYTE;  
 RX MEDLINE=88007472; PubMed=3308864;  
 RA Nakamura T., Hirai T., Tokunaga F., Kawabata S., Iwanaga S.;  
 RT "Purification and amino acid sequence of Kunitz-type protease  
 RT inhibitor found in the hemocytes of horseshoe crab (Tachypleus  
 RT tridentatus).";  
 CC -!- FUNCTION: INHIBITOR OF TRYPSIN AND CHYMOTRYPSIN.  
 CC -!- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.

DR PIR: A26923; A26923.  
 DR HSSP: P00974; 4TPI.  
 DR INTERPRO: IPR002223; -.  
 DR PFAM: PF00014; Kunitz\_BPTI; 1.  
 DR PRINTS: PR00759; BASICPTASE.  
 DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 1.  
 DR PROSITE: PS0279; BPTI\_KUNITZ\_2; 1.  
 KW Serine protease inhibitor.  
 FT DISULFID 8 58 BY SIMILARITY.  
 FT DISULFID 17 41 BY SIMILARITY.  
 FT DISULFID 33 54 BY SIMILARITY.  
 FT ACT\_SITE 18 19 REACTIVE BOND (BY SIMILARITY).  
 SQ SEQUENCE 61 AA; 6825 MW; 730E82CDD0653E48 CRC64;

Query Match 4.1%; Score 7; DB 1; Length 61;  
 Best Local Similarity 100.0%; Pred. No. 1.5;  
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 111 VTGPCRA 117

|||||

Db 13 VTGPCRA 19  
 RESULT 8  
 IVBT\_ERIMA STANDARD; PRT; 62 AA.  
 ID IVBT\_ERIMA  
 AC P24541;  
 DT 01-MAR-1992 (Rel. 21, Created)  
 DT 01-MAR-1992 (Rel. 21, Last sequence update)  
 DT 01-MAY-1992 (Rel. 22, Last annotation update)  
 DE VENOM TRYPSIN INHIBITOR.  
 OS Eristocophis macmahoni (Leaf-nosed viper).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Lepidosauria; Squamata; Scleroglossa; Serpentes; Colubroidea;  
 OC Viperidae; Viperinae; Eristocophis.  
 RN [1]  
 RP SEQUENCE.  
 RC TISSUE=VENOM;  
 RX MEDLINE=92077130; PubMed=1743283;  
 RA Siddiqui A.R., Zaidi Z.H., Joernvall H.;  
 RT "Purification and characterization of a Kunitz-type trypsin inhibitor  
 RT from Leaf-nosed viper venom.";  
 RL FEBS Lett. 294:141-143(1991).  
 CC -!- FUNCTION: THIS PROTEIN INHIBITS TRYPSIN AND KALLIKREIN.  
 CC -!- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 DR PIR: S19327; S19327.  
 DR HSSP: P31713; 1SHP.  
 DR INTERPRO: IPR002223; -.  
 DR PFAM: PF00014; Kunitz\_BPTI; 1.  
 DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 1.  
 DR PROSITE: PS0279; BPTI\_KUNITZ\_2; 1.  
 KW Venom; Serine protease inhibitor.  
 FT DISULFID 2 52 BY SIMILARITY.  
 FT DISULFID 11 35 BY SIMILARITY.  
 FT DISULFID 27 48 BY SIMILARITY.  
 FT ACT\_SITE 12 13 REACTIVE BOND (BY SIMILARITY).  
 SQ SEQUENCE 62 AA; 6772 MW; 0A2ED0ADB20DF938 CRC64;

Query Match 4.1%; Score 7; DB 1; Length 62;  
 Best Local Similarity 100.0%; Pred. No. 1.6;  
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 133 NFYGGC 139

|||||

DB 29 NFYGGC 35

## RESULT 9

NLTP\_DAUCA  
 ID NLTP\_DAUCA STANDARD; PRT; 120 AA.  
 AC P27631;  
 DT 01-AUG-1992 (Rel. 23, Created)  
 DT 01-AUG-1992 (Rel. 23, Last sequence update)  
 DT 01-NOV-1997 (Rel. 35, Last annotation update)  
 DE NONSPECIFIC LIPID-TRANSFER PROTEIN PRECURSOR (LTP) (EXTRACELLULAR  
 DE PROTEIN 2).  
 GN EP2.  
 OS Daucus carota (Carrot).  
 OC Eukaryota; Viridiplantae; Embryophyta; Tracheophyta; Spermatophyta;  
 OC Magnoliophyta; eudicotyledons; core eudicots; Asteridae;  
 OC euasterids II; Apiales; Apiaceae; Daucus.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=CV. NORTHRUP KING;  
 RX MEDLINE=92361243; PubMed=1822991;  
 RA Sterk P., Booldj H., Scheellekens G.A., van Kammen A., de Vries S.C.;  
 RT "Cell-specific expression of the carrot EP2 lipid transfer protein  
 RT gene.";  
 RL Plant Cell 3:907-921(1991).  
 CC -!- FUNCTION: PLANT NONSPECIFIC LIPID-TRANSFER PROTEINS TRANSFER  
 CC PHOSPHOLIPIDS AS WELL AS GALACTOLIPIDS ACROSS MEMBRANES. MAY PLAY  
 CC A ROLE IN WAX OR CUTIN DEPOSITION IN THE CELL WALLS OF EXPANDING

CC EPIDERMAL CELLS AND CERTAIN SECRETORY TISSUES.  
CC -!- TISSUE SPECIFICITY: EXPRESSED IN PROTOPLASM CELLS OF SOMATIC AND  
CC ZYGOTIC EMBRYOS, AND TRANSIENTLY EXPRESSED IN EPIDERMAL CELL  
CC LAYERS OF LEAVES, FLOWERS, AND SEEDS.  
CC -!- SIMILARITY: BELONGS TO THE PLANT LTP FAMILY.  
CC  
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CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).  
CC  
CC EMBL: M64746; AAB96834.1; -  
CC PIR: JQ1280; JQ1280.  
CC HSSP: P19656; IAEH.  
CC INTERPRO: IPR000528; -  
CC PFAM: PF00279; LTP; 1.  
CC PRINTS: PR00382; LIPIDTRNSFR.  
CC PROSITE: PS00597; PLANT\_LTP; 1.  
CC Lipid-binding; Transport; Signal.  
CC SIGNAL 1 26 POTENTIAL.  
CC CHAIN 27 120 NONSPECIFIC LIPID-TRANSFER PROTEIN.  
FT DISULFID 30 79 BY SIMILARITY.  
FT DISULFID 40 56 BY SIMILARITY.  
FT DISULFID 57 102 BY SIMILARITY.  
FT DISULFID 77 116 BY SIMILARITY.  
FT DISULFID 120 120 AA; 12504 MW; E85C6EBA5E592214 CRC64;  
SQ SEQUENCE 120 AA; 12504 MW; E85C6EBA5E592214 CRC64;  
  
Query Match 4.1%; Score 7; DB 1; Length 120;  
Best Local Similarity 100.0%; Pred. No. 2.8;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 107 TANAVTG 113  
Db 83 TANAVTG 89  
  
RESULT 10  
ID IATR\_HORSE STANDARD; PRT; 123 AA.  
AC P04365;  
DT 20-MAR-1987 (Rel. 04, Created)  
DT 20-MAR-1987 (Rel. 04, Last sequence update)  
DT 01-APR-1990 (Rel. 14, Last annotation update)  
DE INTER-ALPHA-TRYPsin INHIBITOR (ITI) (HI-14) (INHIBITORY FRAGMENT OF  
DE ITI) (FRAGMENT).  
OS Equus caballus (Horse).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Perissodactyla; Equidae; Equus.  
RN [1]  
RP MEDLINE:85225967; PubMed:2408637;  
RA Hochstrasser K., Wächter E., Albrecht G.J., Reisinger P.;  
RT "Kunitz-type proteinase inhibitors derived by limited proteolysis of  
RT the inter-alpha-trypsin inhibitor, X. The amino-acid sequences of the  
RT trypsin-released inhibitors from horse and pig inter-alpha-trypsin  
RT inhibitors."  
RL Biol. Chem. Hoppe-Seyler 366:473-478(1985).  
CC  
CC -!- FUNCTION: THIS INHIBITORY FRAGMENT, RELEASED FROM NATIVE ITI AFTER  
CC LIMITED PROTEOLYSIS WITH TRYPSIN, CONTAINS TWO HOMOLOGOUS DOMAINS.  
CC WHEREAS THE SECOND DOMAIN IS A STRONG INHIBITOR OF TRYPSIN, THE  
CC FIRST DOMAIN INTERACTS WEAKLY WITH PMN-GRANULOCYTIC ELASTASE AND  
CC NOT AT ALL WITH PANCREATIC ELASTASE.  
CC -!- MISCELLANEOUS: THE AMINO ACID AT POSITION P2' (17) APPEARS TO  
CC DETERMINE THE SPECIFICITY OF THE INHIBITION OF DOMAIN I.  
CC INHIBITORS WITH METHIONINE IN THIS POSITION INTERACT WEAKLY WITH  
CC CHYMOTRYPSIN AND ELASTASE; THOSE WITH LEUCINE INTERACT STRONGLY.  
CC -!- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
CC PIR: A01210; T1H0B1.  
CC HSSP: P10646; IADZ.

DR INTERPRO: IPR002223; -  
DR PFAM: PF00014; Kunitz\_BPTI; 2.  
DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 2.  
DR PROSITE: PS00279; BPTI\_KUNITZ\_2; 2.  
KW Plasma; Glycoprotein; Serine protease inhibitor; Repeat.  
FT NON\_TER 1 1  
FT DOMAIN 1 56 I.  
FT DOMAIN 57 123 II.  
FT DISULFID 5 55  
FT DISULFID 14 38  
FT DISULFID 30 51  
FT DISULFID 61 111  
FT DISULFID 70 94  
FT DISULFID 86 107  
FT ACT\_SITE 15 16 INHIBITORY SITE (P1) (CHYMOTRYPSIN,  
FT ACT\_SITE 71 72 ELASTASE).  
FT CARBOHYD 24 24 INHIBITORY SITE (P1) (TRYPSIN).  
FT NON\_TER 123 123 N-LINKED (GLCNAC. .).  
SQ SEQUENCE 123 AA; 13510 MW; CE1A9120774411D5 CRC64;  
  
Query Match 4.1%; Score 7; DB 1; Length 123;  
Best Local Similarity 100.0%; Pred. No. 2.9;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 136 YGGCRGN 142  
Db 91 YGGCRGN 97  
  
RESULT 11  
ID TYPE\_BUCAL STANDARD; PRT; 521 AA.  
AC Q44695;  
DT 30-MAY-2000 (Rel. 39, Created)  
DT 30-MAY-2000 (Rel. 39, Last sequence update)  
DT 30-MAY-2000 (Rel. 39, Last annotation update)  
DE ANTHRANILATE SYNTHASE COMPONENT I (EC 4.1.3.27).  
GN TRPE OR BUPT01.  
OS Buchnera aphidicola (subsp. Acyrthosiphon pisum) (Acyrthosiphon pisum  
OS symbiotic bacterium).  
OG Bacteria; Proteobacteria; gamma subdivision; Buchnera.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE:96215866; PubMed:8642610;  
RA Rounbakhsh D., Cai C.-Y., von Dohlen C.D., Clark M.A., Baumann L.,  
RA Baumann P., Moran N.A., Voegtlin D.J.;  
RT "The tryptophan biosynthetic pathway of aphid endosymbionts  
RT (Buchnera): genetics and evolution of plasmid-associated anthranilate  
RT synthase (trpEG) within the aphididae.";  
RL J. Mol. Evol. 42:414-421(1996).  
CC -!- CATALYTIC ACTIVITY: CHORISMATE + L-GLUTAMINE = ANTHRANILATE +  
CC PYRUVATE + L-GLUTAMATE.  
CC -!- PATHWAY: FIRST STEP IN BIOSYNTHESIS OF TRYPTOPHAN  
CC -!- SUBUNIT: TETRAMER OF TWO COMPONENTS I AND TWO COMPONENTS II (B;  
CC -!- MISCELLANEOUS: COMPONENT I CATALYZES THE FORMATION OF ANTHRANILATE  
CC USING AMMONIA RATHER THAN GLUTAMINE, WHEREAS COMPONENT II PROVIDES  
CC GLUTAMINE AMIDOTRANSFERASE ACTIVITY.  
CC -!- SIMILARITY: BELONGS TO THE ANTHRANILATE SYNTHASE COMPONENT I  
CC FAMILY.  
CC  
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CC  
CC EMBL: L43555; AAD09346.1; -

DR PRAM: PF00425; chorismate\_bind; 1.  
DR PRIMS: PRO0095; ANTSNTHASEI.  
KW Tryptophan biosynthesis; Lyase; Plasmid  
SQ SEQUENCE 521 AA; 56655 MW; 285598FE7DF4271 CRC64;

Query Match 4.1%; Score 7; DB 1; Length 521;  
Best Local Similarity 100.0%; Pred. No. 11;  
Matches 7; Conservative 0; Mismatches 0; Gaps 0;  
Indels 0;

QY 12 LVSKVG 18  
| | | | | | | |  
DB 400 LVSKVG 406

RESULT 12  
CO3\_HUMAN  
ID CO3\_HUMAN STANDARD; PRT; 1663 AA.  
AC P01024;  
DT 21-JUL-1986 (Rel. 01, Last sequence update)  
TX 21-JUL-1986 (Rel. 01, Last sequence update)  
FE 15-DEC-1998 (Rel. 37, Last annotation update)  
DE COMPLEMENT C3 PRECURSOR [CONTAINS: C3A ANAPHYLATOXIN].  
GN C3.  
OS Homo sapiens (Human).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
NC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
[1]  
RA "Human anaphylatoxin (C3a) from the third component of complement."  
RX MEDLINE-85140166; PubMed-2579379;  
de Brujin M.H.L.; Fey G.H.;  
"Human complement component C3: cDNA coding sequence and derived  
primary structure.";  
Proc. Natl. Acad. Sci. U.S.A. 82:708-712(1985).  
[2]  
SQ SEQUENCE OF 672-748.  
RX MEDLINE-76069169; PubMed-1238393;  
Hugli T.E.;  
"Human anaphylatoxin (C3a) from the third component of complement."  
Primary structure.";  
J. Biol. Chem. 250:8293-8301(1975).  
[3]  
SQ SEQUENCE OF 1409-1563.  
RX MEDLINE-88154452; PubMed-3279119;  
Daoudaki M.E.; Becherer J.D.; Lambiris J.D.;  
"A 34-amino acid peptide of the third component of complement  
mediates properdin binding.";  
J. Immunol. 140:1577-1580(1988).  
[4]  
SQ SEQUENCE OF 988-1036.  
MEDLINE-82174534; PubMed-6175959;  
Thomas M.L.; Janatova J.; Gray W.R.; Tack B.F.;  
"Third component of human complement: localization of the internal  
thioester bond.";  
Proc. Natl. Acad. Sci. U.S.A. 79:1054-1058(1982).  
[5]  
STRUCTURE BY NMR OF C3A.  
RX MEDLINE-88276894; PubMed-3260670;  
Nettesheim D.G.; Edalji R.P.; Mollison K.W.; Greer J.; Zwieterweg E.R.;  
"Secondary structure of complement component C3a anaphylatoxin in  
solution as determined by NMR spectroscopy: differences between  
crystal and solution conformations.";  
Proc. Natl. Acad. Sci. U.S.A. 85:5036-5040(1988).  
[6]  
MUTAGENESIS OF THIOESTER BOND REGION.  
RX MEDLINE-92250565; PubMed-1577777;  
Isaac L.; Isenman D.E.;  
"Structural requirements for thioester bond formation in human  
complement component C3. Reassessment of the role of thioester bond  
integrity on the conformation of C3.";  
J. Biol. Chem. 267:10062-10069(1992).  
[7]  
DISULFIDE BONDS.

RX MEDLINE-93106233; PubMed-8416818;  
Dolner K.; Sottrup-Jensen L.;  
"Disulfide bridges in human complement component C3b.";  
FEBS Lett. 315:85-90(1993).  
[8]  
RX X-RAY CRYSTALLOGRAPHY (2.0 ANGSTROMS) OF 996-1303.  
MEDLINE-98259089; PubMed-9596584;  
Nagar B.; Jones R.G.; Diefenbach R.J.; Isenman D.E.; Rini J.M.;  
"X-ray crystal structure of C3d: a C3 fragment and ligand for  
complement receptor 2.";  
Science 280:1277-1281(1998).  
[9]  
RX VARIANT C3F/S  
MEDLINE-89309808; PubMed-2473125;  
Pozhansky M.C.; Clissold P.M.; Lachmann P.J.;  
"The difference between human C3f and C3s results from a single amino  
acid change from an asparagine to an aspartate residue at position  
1216 on the alpha-chain of the complement component, C3.";  
J. Immunol. 143:1254-1258(1989).  
[10]  
ERRATUM (RETRACTION OF ABOVE ARTICLE).  
RX MEDLINE-90063087; PubMed-2584723;  
Pozhansky M.C.; Clissold P.M.; Lachmann P.J.;  
J. Immunol. 143:3860-3862(1989).  
[11]  
RX VARIANTS GLY-102 AND PRO-314  
MEDLINE-9101240; PubMed-1976733;  
Botto M.; Yong Fong K.; So A.K.; Koch C.; Walport M.J.;  
"Molecular basis of polymorphisms of human complement component C3.";  
J. Exp. Med. 172:1011-1017(1990).  
[12]  
RX VARIANT ASN-549.  
MEDLINE-95050640; PubMed-7961791;  
Singer L.; Whitehead W.T.; Akama H.; Katz Y.; Fishelson Z.;  
Wetzel R.A.;  
"Inherited human complement C3 deficiency. An amino acid substitution  
in the beta-chain (ASP549 to ASN) impairs C3 secretion.";  
J. Biol. Chem. 269:28494-28499(1994).  
[13]  
RX VARIANT GLN-1320.  
Watanabe Y.; Matsui N.; Yan K.; Nishimukai H.; Tokunaga K.;  
Juji T.; Kobayashi N.; Kohsaka T.;  
"A novel C3 allotype C3\*F02 has an amino acid substitution that may  
inhibit iC3b synthesis and cause C3-hypocomplementemia.";  
Mol. Immunol. 30:62-62(1993).  
CC -!- FUNCTION: C3 PLAYS A CENTRAL ROLE IN THE ACTIVATION OF THE  
COMPLEMENT SYSTEM. ITS PROCESSING BY C3 CONVERTASE IS THE CENTRAL  
REACTION IN BOTH CLASSICAL AND ALTERNATIVE COMPLEMENT PATHWAYS.  
AFTER ACTIVATION C3B CAN BIND COVALENTLY VIA ITS REACTIVE  
THIOLESTER, TO CELL SURFACE CARBOHYDRATES OR IMMUNE AGGREGATES.  
CC -!- FUNCTION: DERIVED FROM PROTEOLYTIC DEGRADATION OF COMPLEMENT C3,  
C3A ANAPHYLATOXIN IS A MEDIATOR OF LOCAL INFLAMMATORY PROCESS. IT  
INDUCES THE CONTRACTION OF SMOOTH MUSCLE, INCREASES VASCULAR  
PERMEABILITY AND CAUSES HISTAMINE RELEASE FROM MAST CELLS AND  
BASOPHILIC LEUKOCYTES.  
CC -!- SUBUNIT: C3 PRECURSOR IS FIRST PROCESSED BY THE REMOVAL OF 4 APC  
RESIDUES, FORMING TWO CHAINS, BETA & ALPHA, LINKED BY A DISULFIDE  
BOND. C3 CONVERTASE ACTIVATES C3 BY CLEAVING THE ALPHA CHAIN,  
RELEASING C3A ANAPHYLATOXIN & GENERATING C3B (BETA CHAIN + ALPHA  
CHAIN).  
CC -!- POLYMORPHISM: THERE ARE TWO ALLELES: C3S (C3 SLOW), THE MOST  
COMMON ALLELE IN ALL RACES AND C3F (C3 FAST), RELATIVELY FREQUENT  
IN CAUCASIIDS, LESS COMMON IN BLACK AMERICAN, EXTREMELY RARE IN  
ORIENTALS.  
CC -!- DISEASE: C3 DEFICIENCY CAUSES A SUSCEPTIBILITY TO PYOGENIC  
INFECTION.  
CC -!- MISCELLANEOUS: C3B IS RAPIDLY SPLIT IN TWO POSITIONS BY FACTOR T  
AND A COFACTOR TO FORM IC3B (INACTIVATED C3B) AND C3F WHICH IS  
RELEASED.  
CC -!- MISCELLANEOUS: IC3B IS THE SLOWLY CLEAVED (POSSIBLY BY FACTOR I)  
TO FORM C3C AND C3D. OTHER PROTEASES PRODUCE OTHER FRAGMENTS SUCH  
AS C3D OR C3G.  
CC -!- SIMILARITY: TO C4, C5 AND ALPHA-2-MACROGLOBULIN.

CC CC -!- SIMILARITY: CONTAINS 1 ANAPHYLATOXIN-LIKE DOMAIN.  
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DR EMBL: K02765; AA85332.1; -  
DR PIR: A01257; C3H0.  
DR PIR: A27603; A27603.  
DR PDB: 1C3D; 18-NOV-98.  
DR SWISS-2DPAGE: P01024; HUMAN.  
DR MIM: 120700; -  
DR INTERPRO: IPR000020; -  
DR INTERPRO: IPR001134; -  
DR INTERPRO: IPR001599; -  
DR INTERPRO: IPR001840; -  
DR INTERPRO: IPR002890; -  
DR PFAM: PF002107; A2M; 1.  
DR PFAM: PF01835; A2M\_LN; 1.  
DR PFAM: PF01821; ANATO; 1.  
DR PFAM: PF01759; NTR; 1.  
DR PRINTS: PRO0004; ANAPHYLATOXN.  
DR PROSITE: PS00477; ALPHA\_2\_MACROGLOBULIN; 1.  
DR PROSITE: PS01177; ANAPHYLATOXIN\_1; 1.  
DR PROSITE: PS01178; ANAPHYLATOXIN\_2; 1.  
DR Complement pathway; Complement alternate pathway; Plasma;  
KW Inflammatory response; Glycoprotein; Signal; Polymorphism;  
KW Disease mutation; 3D-structure.  
FT SIGNAL 1 22  
FT CHAIN 23 1663  
FT CHAIN 23 1663  
FT CHAIN 23 1663  
FT CHAIN 672 1663  
FT PEPTIDE 672 748  
FT CHAIN 749 1663  
FT CHAIN 749 1663  
FT PEPTIDE 749 954  
FT PEPTIDE 955 1303  
FT PEPTIDE 955 1001  
FT PEPTIDE 1002 1303  
FT PEPTIDE 1304 1320  
FT PEPTIDE 1304 1320  
FT SITE 748 749  
FT SITE 954 955  
FT SITE 1303 1304  
FT SITE 1320 1321  
FT DOMAIN 693 728  
FT DOMAIN 1424 1456  
FT DISULFID 559 816  
FT DISULFID 627 662  
FT DISULFID 693 720  
FT DISULFID 694 727  
FT DISULFID 707 728  
FT DISULFID 873 1513  
FT DISULFID 1101 1158  
FT DISULFID 1388 1489  
FT DISULFID 1389 1458  
FT DISULFID 1506 1511  
FT DISULFID 1518 1590  
FT DISULFID 1537 1661  
FT DISULFID 1637 1646  
FT CARBOHYD 85 85  
FT CARBOHYD 939 939  
FT CARBOHYD 1617 1617  
FT THIOLEST 1010 1013  
FT VARIANT 102 102  
FT VARIANT 314 314  
FT VARIANT 549 549  
FT VARIANT 549 549  
Query Match 4.1%; Score 7; DB 1; Length 1663;

Best Local Similarity 100.0%; Pred. No. 31;  
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps  
Oy 46 GNSNNYL 52  
Db 450 GNSNNYL 456  
|||||  
RESULT 13  
LRP2\_RAT  
ID LRP2\_RAT STANDARD; PRT; 4660 AA.  
AC P98158;  
DT 01-OCT-1996 (Rel. 34, Created)  
DT 01-OCT-1996 (Rel. 34, Last sequence update)  
DT 01-OCT-2000 (Rel. 40, Last annotation update)  
DE LOW-DENSITY LIPOPROTEIN RECEPTOR-RELATED PROTEIN 2 PRECURSOR (MEGALIN)  
DE (GLYCOPROTEIN 330).  
GN LRP2.  
OS Rattus norvegicus (Rat).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; R  
[1]  
RN RP SEQUENCE FROM N.A.  
RC STRAIN-SPRAGUE-DAWLEY; TISSUE-KIDNEY;  
RX MEDLINE=95024033; PubMed=7937880;  
RA Saito A., Pietromonaco S., Loo A.K.C., Farquhar M.G.;  
RT "Complete cloning and sequencing of rat gp330/megalin, a  
RT distinctive member of the low density lipoprotein receptor gene  
RT family.";  
RL Proc. Natl. Acad. Sci. U.S.A. 91:9725-9729(1994).  
EN [2]  
FP FUNCTION  
RA MEDLINE=95386696; PubMed=7544804;  
RA Moestrup S.K., Cui S., Vorum H., Bregengaard C., Bjorn S.E.,  
RA Norris K., Gliemann J., Christensen E.I.;  
RT "Evidence that epithelial glycoprotein 330/megalin mediates uptake of  
RT polybasic drugs.";  
RL J. Clin. Invest. 96:1404-1413(1995).  
EN [3]  
FP TISSUE SPECIFICITY.  
RA MEDLINE=94172242; PubMed=7510321;  
RA Zheng G., Bachinsky D.R., Stamenkovic I., Strickland D.K., Brown D.,  
RA "Organ distribution in rats of two members of the low-density  
RT lipoprotein receptor gene family, gp330 and LRP/alpha 2MR, and the  
RT receptor-associated protein (RAP).";  
RL J. Histochem. Cytochem. 42:531-542(1994).  
CC -!- FUNCTION: BINDS PLASMINOGEN, EXTRACELLULAR MATRIX COMPONENTS,  
CC PLASMINOGEN ACTIVATOR-PLASMINOGEN ACTIVATOR INHIBITOR TYPE I  
CC COMPLEX, APOLIPOPROTEIN E-ENRICHED BETA-VLDL, LIPOPROTEIN LIPASE,  
CC LACTOFERRIN; CLUSTERIN AND CALCIUM.  
CC -!- FUNCTION: RECEPTOR-MEDIATED UPTAKE OF POLYBASIC DRUGS SUCH AS  
CC APROTININ, AMINOGLYCOSIDES AND POLYMYXIN B.  
CC -!- SUBUNIT: FORMS A MULTIMERIC COMPLEX TOGETHER WITH A RECEPTOR-  
CC ASSOCIATED PROTEIN (RAP).  
CC -!- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN. EXPRESSED IN  
CC CLATHRIN-COATED PITS; A SOLUBLE FORM IS POSSIBLY DERIVED BY  
CC CLEAVAGE AT THE CELL SURFACE.  
CC -!- TISSUE SPECIFICITY: EPITHELIAL CELLS OF KIDNEY GLOMERULUS AND  
CC PROXIMAL TUBULE, LUNG, EPIDIDYMIS, YOLK SAC, AMONG OTHERS.  
CC -!- SIMILARITY: CONTAINS 36 LDL-RECEPTOR CLASS A DOMAINS.  
CC -!- SIMILARITY: CONTAINS 37 LDL-RECEPTOR CLASS B DOMAINS.  
CC -!- SIMILARITY: CONTAINS 17 EGF-LIKE DOMAINS.  
CC -!- SIMILARITY: CONTAINS 3 SH3-BINDING DOMAINS.  
CC -!- SIMILARITY: CONTAINS 1 SH2-BINDING DOMAIN.  
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CC EMBL: L34049; AAA51369.1; -  
DR HSSP; P01130; 1AJJ.  
DR INTERPRO; IPR000033; -  
DR INTERPRO; IPR000152; -  
DR INTERPRO; IPR000561; -  
DR INTERPRO; IPR001881; -  
DR INTERPRO; IPR002172; -  
DR PFAM; PF00057; ldl\_recept\_a; 36.  
DR PFAM; PF00058; ldl\_recept\_b; 33.  
DR PRINTS; PR0261; LDLRECEPTOR.  
DR PROSITE; PS00010; ASX\_HYDROXIL; 4.  
DR PROSITE; PS00022; EGF\_1; 1.  
DR PROSITE; PS01186; EGF\_2; 8.  
DR PROSITE; PS01187; EGF\_CA; 3.  
DR PROSITE; PS01209; LDLRA\_1; 31.  
DR PROSITE; PS00068; LDLRA\_2; 36.  
KW Glycoprotein; Repeat; Endocytosis; Coated pits; Transmembrane;  
KW Receptor; EGF-like domain; Signal.  
FT CHAIN 1 25  
FT SIGNAL 1 25  
FT CHAIN 26 4660  
FT DOMAIN 26 4425  
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RESULT 14  
ISF2\_GALME STANDARD; PRT; 52 AA.  
AC P81906;  
DT 30-MAY-2000 (Rel. 39, Created)  
DT 30-MAY-2000 (Rel. 39, Last sequence update)  
DT 30-MAY-2000 (Rel. 39, Last annotation update)  
DE INDUCIBLE SERINE PROTEASE INHIBITOR 2 (ISPI-2) (FRAGMENT).  
OS Galleria mellonella (Wax moth).  
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;  
OC Pterygota; Neoptera; Endopterygota; Lepidoptera; Glossata; Ditrysia;  
OC Pyraloidea; Pyralidae; Galleriinae; Galleria.  
RN [1]  
RP SEQUENCE.  
RX TISSUE=HEMOLYMPH;  
RA MEDLINE=20193629; PubMed=10727944;  
RA Froebius A.C., Kanost M.R., Goetz P., Vilcinskis A.;  
RA "Isolation and characterization of novel inducible serine protease  
inhibitors from larval hemolymph of the greater wax moth Galleria  
mellonella";  
RL Eur. J. Biochem. 267:2046-2053(2000).  
CC -!- FUNCTION: INHIBITS TRYPSIN AND THE TOXIN PROTEASE PR2 OF M.  
CC ANISOPLIAE. DOES NOT INHIBIT CHYMOTRYPSIN, SUBTILISIN CARLSBERG,  
CC PROTEINASE K, PORCINE PANCREATIC ELASTASE AND THE TOXIN PROTEASE  
CC PR1 OF M. ANISOPLIAE.  
CC -!- DEVELOPMENTAL STAGE: LAST INSTAR LARVAE.  
CC -!- INDUCTION: BY INFECTION.  
CC -!- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
DR INTERPRO: IPR002223; -.  
DR PFAM: PF00014; Kunitz\_BPTI; 1.  
DR PRINTS: PR00759; BASICPTASE.  
DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 1.  
-DR PROSITE: PS50279; BPTI\_KUNITZ\_2; 1.  
KW Serine protease inhibitor.  
FT DISULFID 14 38 BY SIMILARITY.  
FT ACT\_SITE 30 51 BY SIMILARITY.  
FT NON\_TER 15 16 REACTIVE BOND (BY SIMILARITY).  
FT SEQUENCE 52 AA; 6057 MW; 31CED34D59C42ABE CRC64;

QY 39 FVYGGC 44  
Db 33 FVYGGC 38  
Query Match 3.5%; Score 6; DB 1; Length 52;  
Best Local Similarity 100.0%; Pred. No. 15;  
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 15  
ISH1\_STOHE STANDARD; PRT; 55 AA.  
AC P31713;  
DT 01-JUL-1993 (Rel. 26, Created)  
DT 01-JUL-1993 (Rel. 26, Last sequence update)  
DT 15-JUL-1998 (Rel. 36, Last annotation update)  
DE KUNITZ-TYPE PROTEINASE INHIBITOR SHPI-1  
OS Stoichactis helianthus (Caribbean sea anemone) (Stichodactyla  
helianthus).  
OC Eukaryota; Metazoa; Cnidaria; Anthozoa; Zoantharia; Actiniaria;  
OC Stichodactylidae; Stichodactyla.  
RN [1]  
RP SEQUENCE.  
RX MEDLINE=97179757; PubMed=9027993;  
RA Delfin J., Martinez J., Antuch W., Morera V., Gonzalez Y.,  
RA Rodriguez R., Marquez M., Saroyan A., Larionova N., Diaz J.,  
RA Padron G., Chavez M.;  
RA "Purification, characterization and immobilization of proteinase  
inhibitors from Stichodactyla helianthus";  
RL Toxicon 34:1367-1376(1996).  
RN [2]

RP STRUCTURE BY NMR, AND DISULFIDE BONDS.  
RX MEDLINE=93215644; PubMed=8462542;  
RA Antuch W., Berndt K.D., Chavez M.A., Delfin J., Wuethrich K.;  
RA "The NMR solution structure of a Kunitz-type proteinase inhibitor  
from the sea anemone Stichodactyla helianthus";  
RL Eur. J. Biochem. 212:675-684(1993).  
CC -!- FUNCTION: ACTIVE AGAINST SERINE, CYSTEINE, AND ASPARTIC  
CC PROTEINASES.  
CC -!- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
DR PIR: S30332; S30332.  
DR PDB: 1SHP; 31-JAN-94.  
DR INTERPRO: IPR002223; -.  
DR PFAM: PF00014; Kunitz\_BPTI; 1.  
DR PROSITE: PS00280; BPTI\_KUNITZ\_1; 1.  
DR PROSITE: PS50279; BPTI\_KUNITZ\_2; 1.  
KW Serine protease inhibitor; 3D-structure.  
FT DISULFID 3 53  
FT ACT\_SITE 12 36  
FT DISULFID 28 49  
FT ACT\_SITE 13 14  
FT HELIX 2 4  
FT STRAND 17 22  
FT TURN 23 26  
FT STRAND 27 32  
FT TURN 36 37  
FT STRAND 43 43  
FT HELIX 45 53  
SQ SEQUENCE 55 AA; 6116 MW; 532B96E3127000D4 CRC64;

Query Match 3.5%; Score 6; DB 1; Length 55;  
Best Local Similarity 100.0%; Pred. No. 16;  
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 FVYGGC 139  
Db 31 FVYGGC 36

Search completed: January 31, 2001, 15:07:48  
Job time: 116 sec

Wed Jan 31 15:14:33 2001

us-09-441-654a-1.oli.rsp

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GenCore version 4.5  
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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:05:23 ; Search time 22.96 Seconds  
(without alignments)  
867.829 Million cell updates/sec

Title: US-09-441-654A-1  
Perfect score: 170  
Sequence: 1 ADERSIHDFCLVSKVVGRC.....ACMLRCFRQENPPLPLGSK 170

Scoring table: OLIGO  
Gapop 60.0 , Gapext 60.0

Searched: 374700 seqs, 117207915 residues  
Hit size: 0

Total number of hits satisfying chosen parameters: 374700

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

- Database : SPTREMBL15.\*
- 1: sp\_archaea.\*
  - 2: sp\_bacteria.\*
  - 3: sp\_fungi.\*
  - 4: sp\_human.\*
  - 5: sp\_invertebrate.\*
  - 6: sp\_mammal.\*
  - 7: sp\_mhc.\*
  - 8: sp\_organelle.\*
  - 9: sp\_phase.\*
  - 10: sp\_plant.\*
  - 11: sp\_rodent.\*
  - 12: sp\_virus.\*
  - 13: sp\_vertebrate.\*
  - 14: sp\_unclassified.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	170	100.0	252	4	O43291
2	170	100.0	252	4	O00271
3	144	84.7	252	4	O14895
4	13	7.6	195	11	O9WU04
5	13	7.6	252	11	O9WU03
6	9	5.3	58	5	O9TWG0
7	9	5.3	58	5	O9TWG9
8	8	4.7	59	5	O9TWG8
9	8	4.7	922	5	O21418
10	8	4.7	1195	5	O9N343
11	8	4.7	1599	5	O09983
12	7	4.1	110	5	O9VQU0
13	7	4.1	183	5	O16784
14	7	4.1	210	13	O91443
15	7	4.1	230	13	O13000
16	7	4.1	253	2	O54224
17	7	4.1	260	5	O46164
18	7	4.1	355	5	O9VQ08
19	7	4.1	438	4	O9Y4N9

20	7	4.1	499	2	Q9L2G7	Q9L2G7 streptomyce
21	7	4.1	500	2	Q9ZN02	Q9ZN02 helicobacte
22	7	4.1	507	2	O87080	O87080 vibrio chol
23	7	4.1	514	5	O9TWE5	O9TWE5 carcinoscor
24	7	4.1	521	2	O9KGQ2	O9KGQ2 buchnera sp
25	7	4.1	521	2	O9W3R0	O9W3R0 drosophila
26	7	4.1	654	5	O9VFX6	O9VFX6 drosophila
27	7	4.1	719	5	O9U021	O9U021 giardia lam
28	7	4.1	719	5	O9U019	O9U019 giardia lam
29	7	4.1	763	5	O9XZD0	O9XZD0 drosophila
30	7	4.1	836	4	O94856	O94856 homo sapien
31	7	4.1	972	5	O44938	O44938 haemochus
32	7	4.1	974	13	Q91735	Q91735 xenopus lae
33	7	4.1	988	13	Q07498	Q07498 gallus gall
34	7	4.1	1151	11	Q9QVNS	Q9QVNS rattus sp.
35	7	4.1	1214	5	Q25338	Q25338 latrodectus
36	7	4.1	1474	5	O62504	O62504 caenorhabdi
37	7	4.1	1668	4	O15026	O15026 homo sapien
38	7	4.1	1787	10	O9S7P0	O9S7P0 arabidopsis
39	7	4.1	2167	5	O76840	O76840 caenorhabdi
40	7	4.1	2208	5	Q09515	Q09515 caenorhabdi
41	7	4.1	2230	5	Q9NAV4	Q9NAV4 drosophila
42	7	4.1	2971	4	Q9Y5L9	Q9Y5L9 homo sapien
43	7	4.1	3198	5	Q9U8G8	Q9U8G8 mangusta
44	6	3.5	41	12	O11552	O11552 mus muscu
45	6	3.5	42	2	Q49201	Q49201 myocorinae

ALIGNMENTS

RESULT 1

O43291

ID O43291 PRELIMINARY; PRT; 252 AA.

AC O43291;

DT 01-JUN-1998 (TREMBLrel. 06, Created)

DT 01-JUN-1998 (TREMBLrel. 06, Last sequence update)

DE 01-OCT-2000 (TREMBLrel. 15, Last annotation update)

DE HEPATOCYTE GROWTH FACTOR ACTIVATOR INHIBITOR TYPE 2.

OS Homo sapiens (Human)

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

OX NCBI\_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=98010584; PubMed=9346890;

RA Kawaguchi T., Qin L., Shimomura T., Kondo J., Matsumoto K., Denda K.,

RA Kitamura N.;

RT "Purification and cloning of hepatocyte growth factor activator

RT inhibitor type 2, a Kunitz-type serine protease inhibitor.";

RL J. Biol. Chem. 272:27558-27564(1997).

DR EMBL: AB006534; BAA25024.1; -.

DR HSSP: P05067; ITAW.

DR INTERPRO: IPR002223; -.

DR PFAM: PF00014; Kunitz\_BPTI; 2.

DR PRINTS: PR00759; BASICPTASE.

DR PROSITE: PS00280; BPTI\_KUNITZ; 2.

KW Serine protease inhibitor.

SQ SEQUENCE 252 AA; 28189 MW; F7D3D834ED631DF0 CRC64;

Query Match 100.0%; Score 170; DB 4; Length 252;  
Best Local Similarity 100.0%; Pred. No. 1.1e-178;  
Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ADERSIHDFCLVSKVVGRCRASMFRWTVNTDGGSCQLFVYGGCDGNSNNYLKKECLKK 60  
|||||  
Db 28 ADERSIHDFCLVSKVVGRCRASMFRWTVNTDGGSCQLFVYGGCDGNSNNYLKKECLKK 87  
Qy 61 CATVTENATGDLATSRNAADSSVPSAPRQDSEHSSDMFNVEEYCTANAVTGPCRASEP 120  
|||||  
Db 88 CATVTENATGDLATSRNAADSSVPSAPRQDSEHSSDMFNVEEYCTANAVTGPCRASEP 147

QY	121	RWYFDVERNSCNFIYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK	170
Db	148	RWYFDVERNSCNFIYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK	197
RESULT	2		
ID	000271	PRELIMINARY;	PRT; 252 AA.
AC	000271;		
DT	01-JUN-1997	(TReMBLrel. 04, Created)	
DT	01-JUL-1997	(TReMBLrel. 04, Last sequence update)	
DT	01-OCT-2000	(TReMBLrel. 15, Last annotation update)	
DE	BKUNIN.		
OS	Homo sapiens (Human).		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
OX	NCBI_TaxID=9606;		
RN	[1]		
RP	SEQUENCE FROM N.A.		
RC	TISSUE=PLACENTA;		
RA	MEDLINE=97277372; PubMed=9115294;		
RA	Marlor C.W., Delaria K.A., Davis G., Muller D.K., Greve J.M.,		
RA	Tamburini P.P.;		
RT	"Identification and cloning of human placental bikunin, a novel serine		
RT	protease inhibitor containing two Kunitz domains.";		
RL	J. Biol. Chem. 272:12202-12208(1997).		
DR	EMBL; U78095; AAC02781.1; -;		
DR	HSSP; P05067; ITAW.		
DR	INTERPRO; IPR002223; -;		
DR	PFAM; PF00014; Kunitz_BPTI; 2.		
DR	PRINTS; PR00759; BASICTPASE.		
DR	PROSITE; PS00280; BPTI_KUNITZ; 2.		
KW	Serine protease inhibitor.		
SQ	SEQUENCE 252 AA; 28228 MW; A7D3360C0EECAB2B CRC64;		
Query Match	100.0%;	Score 170;	DB 4; Length 252;
Best Local Similarity	100.0%;	Pred. No. 1.1e-178;	
Matches 170;	Conservative 0;	Mismatches 0;	Indels 0; Gaps 0;
QY	1	ADRSRSHDFCLVSKVYVGRASMPRWYNTDGSQCLFYGGCDGNSNNYLTKKECLKK	60
Db	28	ADRSRSHDFCLVSKVYVGRASMPRWYNTDGSQCLFYGGCDGNSNNYLTKKECLKK	87
QY	61	CATVTENATGDLATSNAAADSSVPSAPRODSEHSDMFNYEYCTANAVTGPCRASFP	120
Db	88	CATVTENATGDLATSNAAADSSVPSAPRODSEHSDMFNYEYCTANAVTGPCRASFP	147
121	RWYFDVERNSCNFIYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK	170	
148	RWYFDVERNSCNFIYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK	197	
RESULT	3		
ID	014895	PRELIMINARY;	PRT; 252 AA.
AC	014895;		
DT	01-JAN-1998	(TReMBLrel. 05, Created)	
DT	01-JAN-1998	(TReMBLrel. 05, Last sequence update)	
DT	01-OCT-2000	(TReMBLrel. 15, Last annotation update)	
DE	KUNITZ-TYPE PROTEASE INHIBITOR.		
GN	KOP.		
OS	Homo sapiens (Human).		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
OX	NCBI_TaxID=9606;		
RN	[1]		
RP	SEQUENCE FROM N.A.		
RC	TISSUE=PANCREATIC CANCER;		
RA	Mueller-Pillasch F., Wallrapp C., Bartels K., Varga G., Friess H.,		
RA	Buechler M., Adler G., Gress T.M.;		
RL	Biochim. Biophys. Acta 0:0-0(1997).		
DR	EMBL; AF027205; AAB84031.1; -;		
Query Match	100.0%;	Score 13;	DB 11; Length 195;
Best Local Similarity	100.0%;	Pred. No. 3.5e-06;	
Matches 13;	Conservative 0;	Mismatches 0;	Indels 0; Gaps 0;
QY	134	FIYGGCRGNKNSY	146
Db	104	FIYGGCRGNKNSY	116
RESULT	5		
ID	Q9WU03	PRELIMINARY;	PRT; 252 AA.
AC	Q9WU03;		
DT	01-NOV-1999	(TReMBLrel. 12, Created)	
DT	01-NOV-1999	(TReMBLrel. 12, Last sequence update)	

01-OCT-2000 (TReMBLrel. 15, Last annotation update)  
DE HEPATOCYTE GROWTH FACTOR ACTIVATOR INHIBITOR TYPE 2.  
GN HAI2.  
OS Mus musculus (Mouse).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
OX NCBI\_TaxID=10090;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=BALB/C.  
RX MEDLINE=99160423; PubMed=10049781;  
RA Itoh H., Kataoka H., Hamasuna R., Kitamura N., Koono M.;  
RT "Hepatocyte growth factor activator inhibitor type 2 lacking the first  
Kunitz-type serine proteinase inhibitor domain is a predominant  
product in mouse but not in human."  
RL Biochem. Biophys. Res. Commun. 255:740-748(1999).  
DR EMBL; AF099016; RAD2172.1; -.  
DR HSSP; P05067; ITAW.  
DR INTERPRO; IPR002223; -.  
DR PFAM; PF00014; Kunitz\_BPTI; 2.  
DR PRINTS; PR00759; BASICTPASE.  
DR PROSITE; PS00280; BPTI\_KUNITZ; 2.  
KW Serine protease inhibitor.  
SQ SEQUENCE 252 AA; 27914 MW; B2FF4B86924D4F8F CRC64;

Query Match 7.6%; Score 13; DB 11; Length 252;  
Best Local Similarity 100.0%; Pred. No. 4.4e-06;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 FIYGGCRGNKNSY 146  
Db 161 FIYGGCRGNKNSY 173  
|||||

RESULT 6  
Q9TWF8  
ID Q9TWF8 PRELIMINARY; PRT; 58 AA.  
AC Q9TWF8  
DT 01-MAY-2000 (TReMBLrel. 13, Created)  
DT 01-MAY-2000 (TReMBLrel. 13, Last sequence update)  
DE KALICLUDINE 1, ASKC1.  
DE KALICLUDINE 1, ASKC1.  
OS Anemonia sulcata (Snake-locks sea anemone).  
OC Eukaryota; Metazoa; Chidaria; Anthozoa; Zoantharia; Actiniaria;  
OC Nynantheae; Actiniidae; Anemonia.  
OX NCBI\_TaxID=6108;  
RN [1]  
RP SEQUENCE.  
RX MEDLINE=96027617; PubMed=7559645;  
RA Schweitz H., Bruhn T., Guillemare E., Moinier D., Lancelin J.M.,  
RA Beress L., Lazdunski M.;  
RT "Kalicludines and kaliseptine. Two different classes of sea anemone  
toxins for voltage sensitive K+ channels."  
RL J. Biol. Chem. 270:25121-25126(1995).  
DR HSSP; P10646; 1ADZ.  
DR INTERPRO; IPR002223; -.  
DR PFAM; PF00014; Kunitz\_BPTI; 1.  
DR PRINTS; PR00759; BASICTPASE.  
DR PROSITE; PS00280; BPTI\_KUNITZ; 1.  
SQ SEQUENCE 58 AA; 6691 MW; 253E068896B4BDCD CRC64;

Query Match 5.3%; Score 9; DB 5; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.031;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 FIYGGCRGN 142  
Db 33 FIYGGCRGN 41  
|||||

RESULT 7

Q9TWF9  
ID Q9TWF9 PRELIMINARY; PRT; 58 AA.  
AC Q9TWF9  
DT 01-MAY-2000 (TReMBLrel. 13, Created)  
DT 01-MAY-2000 (TReMBLrel. 13, Last sequence update)  
DE KALICLUDINE 2, ASKC2.  
DE KALICLUDINE 2, ASKC2.  
OS Anemonia sulcata (Snake-locks sea anemone).  
OC Eukaryota; Metazoa; Chidaria; Anthozoa; Zoantharia; Actiniaria;  
OC Nynantheae; Actiniidae; Anemonia.  
OX NCBI\_TaxID=6108;  
RN [1]  
RP SEQUENCE.  
RX MEDLINE=96027617; PubMed=7559645;  
RA Schweitz H., Bruhn T., Guillemare E., Moinier D., Lancelin J.M.,  
RA Beress L., Lazdunski M.;  
RT "Kalicludines and kaliseptine. Two different classes of sea anemone  
toxins for voltage sensitive K+ channels."  
RL J. Biol. Chem. 270:25121-25126(1995).  
DR HSSP; P12111; 2KNT.  
DR INTERPRO; IPR002223; -.  
DR PFAM; PF00014; Kunitz\_BPTI; 1.  
DR PRINTS; PR00759; BASICTPASE.  
DR PROSITE; PS00280; BPTI\_KUNITZ; 1.  
SQ SEQUENCE 58 AA; 6778 MW; F102E71682F1A55C CRC64;

Query Match 5.3%; Score 9; DB 5; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.031;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 FIYGGCRGN 142  
Db 33 FIYGGCRGN 41  
|||||

RESULT 8  
Q9TWF8  
ID Q9TWF8 PRELIMINARY; PRT; 59 AA.  
AC Q9TWF8  
DT 01-MAY-2000 (TReMBLrel. 13, Created)  
DT 01-MAY-2000 (TReMBLrel. 13, Last sequence update)  
DE KALICLUDINE 3, ASKC3.  
DE KALICLUDINE 3, ASKC3.  
OS Anemonia sulcata (Snake-locks sea anemone).  
OC Eukaryota; Metazoa; Chidaria; Anthozoa; Zoantharia; Actiniaria;  
OC Nynantheae; Actiniidae; Anemonia.  
OX NCBI\_TaxID=6108;  
RN [1]  
RP SEQUENCE.  
RX MEDLINE=96027617; PubMed=7559645;  
RA Schweitz H., Bruhn T., Guillemare E., Moinier D., Lancelin J.M.,  
RA Beress L., Lazdunski M.;  
RT "Kalicludines and kaliseptine. Two different classes of sea anemone  
toxins for voltage sensitive K+ channels."  
RL J. Biol. Chem. 270:25121-25126(1995).  
DR HSSP; P31713; 1SHP.  
DR INTERPRO; IPR002223; -.  
DR PFAM; PF00014; Kunitz\_BPTI; 1.  
DR PRINTS; PR00759; BASICTPASE.  
DR PROSITE; PS00280; BPTI\_KUNITZ; 1.  
SQ SEQUENCE 59 AA; 6738 MW; 0C7695C3F394D4A5 CRC64;

Query Match 4.7%; Score 8; DB 5; Length 59;  
Best Local Similarity 100.0%; Pred. No. 0.4;  
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 KVVGCRGA 22  
Db 9 KVVGCRGA 16  
|||||

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RT investigating biology. The C. elegans Sequencing Consortium."
RL Science 282:2012-2018(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-BRISTOL N2;
RA Waterston R.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC024830; AAF59608.1; -
SQ SEQUENCE 1195 AA; 131342 MW; E77C3A6DF2272A18 CRC64;

Query Match 4.7%; Score 8; DB 5; Length 1195;
Best Local Similarity 100.0%; Pred. No. 5.3;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 GCDGNSNN 50
DB 887 GCDGNSNN 894

RESULT 11
Q09983 PRELIMINARY; PRT; 1599 AA.
ID Q09983;
AC Q09983;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-OCT-2000 (TrEMBLrel. 15, Last annotation update)
DE HYPOTHEICAL 171.7 KDA PROTEIN F30H5.3 IN CHROMOSOME III.
GN F30H5.3.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-BRISTOL N2;
RA Pauley A., Steeves L.;
RL Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: STRONG, TO C.ELEGANS ZC84.1.
CC -1- SIMILARITY: BELONGS TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.
DR EMBL; U29096; AAA68408.1; -
DR HSP; P10646; 1TFX.
DR WORMPEP; F30H5.3; CE01927.
DR INTERPRO; IPR002198; -
DR INTERPRO; IPR002223; -
DR INTERPRO; IPR002899; -
DR PFAM; PF01683; EB; 3.
DR PRINTS; PR00759; BASICPTASE.
DR PROSITE; PS00061; ADH_SHORT; UNKNOWN_1.
DR PROSITE; PS00280; BPTI_KUNITZ; 2.
KW Hypothetical protein; Serine protease inhibitor.
FT DOMAIN 7. 10 POLY-LEU.
FT DOMAIN 1520 1523 POLY-GLU.
SQ SEQUENCE 1599 AA; 171658 MW; AB5E6A1D86E9880D CRC64;

Query Match 4.7%; Score 8; DB 5; Length 1599;
Best Local Similarity 100.0%; Pred. No. 6.8;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 GCDGNSNN 50
DB 704 GCDGNSNN 711

RESULT 12
Q09983 PRELIMINARY; PRT; 110 AA.
ID Q09983;
AC Q09983;
DT 01-MAY-2000 (TrEMBLrel. 13, Created)
DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DT 01-OCT-2000 (TrEMBLrel. 15, Last annotation update)
RT "Genome sequence of the nematode C. elegans: a platform for
```

DE CG10031 PROTEIN.  
 GN CG10031.  
 OS Drosophila melanogaster (Fruit fly).  
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Insecta;  
 CC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
 OC Nephelidae; Drosophilidae; Drosophila.  
 NCBI\_TaxID=7227;  
 [1]  
 RN  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=BERKELEY;  
 RX MEDLINE=20196006; PubMed=10731132;  
 RA Adams M.D., Celnik S.E., Holt R.A., Evans C.A., Gocayne J.D.,  
 RA Amanatides P.G., Scher S.E., Li P.W., Hoskins R.A., Galie R.F.,  
 RA George R.A., Lewis S.J., Richards S., Ashburner M., Henderson S.N.,  
 RA Sutton G.G., Wortman J.R., Randell M.D., Zhang Q., Chen L.X.,  
 RA Brandon R.C., Rogers Y.-H.C., Blazek R.G., Champe M., Pfeiffer B.D.,  
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfankuch C., Baldwin D.,  
 RA Ballou R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,  
 RA Beeson K.V., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,  
 RA Borkova D., Botchan M.R., Bouch J., Brokstein P., Brottier P.,  
 RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,  
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,  
 RA de Pablo J.B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,  
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,  
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,  
 RA Folsler C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,  
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,  
 RA Hostin D., Houston K.A., Howland T.J., Ke Z., Kennison J.A., Ketchum K.A.,  
 RA Jalali M., Kalush F., Karpen G.H., Ji J., Wei M.-H., Ibegwam C.,  
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,  
 RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,  
 RA Merkulov G., Milshina N.V., Moberg B., Murphy L., Muzny D.M., Nelson D.L.,  
 RA Mount D.R., Nelson K.A., Nixon K., Nuskern D.R., Pacleb J.M.,  
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,  
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,  
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,  
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,  
 RA Svrtkar R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,  
 RA Wang Z.-Y., Wassarman D.A., Weinstock G.M., Weissbach J.,  
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,  
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,  
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,  
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;  
 "The genome sequence of Drosophila melanogaster.";  
 Science 287:2185-2193(2000).  
 EMBL; AF003579; AAF51074.1; -;  
 HSSP; P12111; 2KNT.  
 DR FLXBASE; FBgn0031563; CG10031.  
 DR INTERPRO; IPR002223; -;  
 DR PFAM; PF00014; Kunitz\_BPTI; 1.  
 DR PRINTS; PR00759; BASICPRASE.  
 SQ SEQUENCE 110 AA; 12240 MW; BB3F2DF4A7EF509D CRC64;

Query Match 4.1%; Score 7; DB 5; Length 110;  
 Best Local Similarity 100.0%; Pred. No. 8.5;  
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 136 YGGCRGN 142  
 Db 86 YGGCRGN 92  
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RESULT 13  
 OL6784  
 ID OL6784 PRELIMINARY; PRT; 183 AA.  
 AC OL6784;  
 DT 01-JAN-1998 (TrEMBLrel. 05, Created)  
 DR EMBL; Z50082; CAA90413.1; -;  
 DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)

DT 01-OCT-2000 (TrEMBLrel. 15, Last annotation update)  
 DE SIMILAR TO THE BPTI/KUNITZ FAMILY OF INHIBITORS.  
 GN T21D12.12.  
 OS Caenorhabditis elegans.  
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;  
 CC Rhabditidae; Peloderinae; Caenorhabditis.  
 NCBI\_TaxID=6239;  
 [1]  
 RN  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=BRISTOL N2;  
 RX MEDLINE=94150718; PubMed=7906398;  
 RA Willson R., Ainscough R., Anderson K., Baynes C., Berks M.,  
 RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,  
 RA Craxton M., Dear S., Du Z., Durbin R., Favello A., Fulton L.,  
 RA Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,  
 RA Jones M., Kershaw J., Kirsten J., Laister N., Latreille P.,  
 RA Lightning J., Lloyd C., McMurray A., Mortimore B., O'Callaghan M.,  
 RA Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Showkeen R.,  
 RA Smaildon N., Smith A., Sonhammer E., Staden R., Sulston J.,  
 RA Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K., Waterston R.,  
 RA Watson A., Weinstock L., Wilkinson-Sproat J., Wohlman P.,  
 RA "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.  
 elegans.";  
 RL Nature 368:32-38(1994).  
 [2]  
 RN  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=BRISTOL N2;  
 RX Woessner J.;  
 RL Submitted (AUG-1997) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AF016687; AAC48097.1; -;  
 DR HSSP; P10646; 1TFX.  
 DR INTERPRO; IPR002223; -;  
 DR PFAM; PF00014; Kunitz\_BPTI; 2.  
 DR PROSITE; PS00280; BPTI\_KUNITZ; 1.  
 KW Serine protease inhibitor.  
 SQ SEQUENCE 183 AA; 20143 MW; CCBE4BE2293CE32A CRC64;

Query Match 4.1%; Score 7; DB 5; Length 183;  
 Best Local Similarity 100.0%; Pred. No. 13;  
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 43 GCDGNSN 49  
 Db 58 GCDGNSN 64  
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RESULT 14  
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 ID Q91443 PRELIMINARY; PRT; 210 AA.  
 AC Q91443;  
 DT 01-NOV-1996 (TrEMBLrel. 01, Created)  
 DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)  
 DE 01-OCT-2000 (TrEMBLrel. 15, Last annotation update)  
 DE INSULIN-LIKE GROWTH FACTOR II PRECURSOR.  
 OS Squalus acanthias (Spiny dogfish).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Chondrichthyes;  
 CC Elasmobranchii; Squala; Squalidae; Squalus.  
 NCBI\_TaxID=7797;  
 [1]  
 RN  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=LIVER;  
 RX MEDLINE=95394151; PubMed=7545136;  
 RA Duquay S.J., Chan S.J., Mommens T.P., Steiner D.F.;  
 RT "Divergence of insulin-like growth factors I and II in the  
 elasmobranch, Squalus acanthias.";  
 RL FEBS Lett. 371:59-72(1995).  
 CC -!- SUBCELLULAR LOCATION: SECRETED (BY SIMILARITY).  
 CC -!- SIMILARITY: BELONGS TO THE INSULIN/IGF/RELAXIN FAMILY.  
 DR EMBL; Z50082; CAA90413.1; -;  
 DR HSSP; P01344; 1GF2.  
 DR INTERPRO; IPR000739; -;  
 DR PFAM; PF00049; Insulin; 1.

us-09-441-654a-1.oli.rspt

Wed Jan 31 15:14:34 2001

DR PRINTS: PR00276; INSULINA.  
 DR PRINTS: PR00277; INSULINB.  
 DR PROSITE: PS00262; INSULIN; 1.  
 DR PRODOM: PD001048; -; 1.  
 KW Signal. 1 49 POTENTIAL.  
 FT SIGNAL 50 210 INSULIN-LIKE GROWTH FACTOR II.  
 FT CHAIN 50 210  
 FT SEQUENCE 210 AA; 23027 MW; 9B433B7C4749A03A CRC64;

Query Match 4.1%; Score 7; DB 13; Length 210;  
 Best Local Similarity 100.0%; Pred.No.15;  
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 VSKVGR 19  
 DB 78 VSKVGR 84

RESULT 15  
 3000 PRELIMINARY; PRT: 230 AA.  
 AC 013000;  
 DT 01-JUL-1997 (TREMblrel. 04, Created)  
 DT 01-JUL-1997 (TREMblrel. 04, Last sequence update)  
 DT 01-OCT-2000 (TREMblrel. 15, Last annotation update)  
 DE PROTEASOME SUBUNIT Y (EC 3.4.99.46).  
 OS Xenopus laevis (African clawed frog).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;  
 OC Xenopodinae; Xenopus.  
 OC NCBI\_TaxID=8355;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=LIVER;  
 RA Nonaka M., Namikawa C., Sasaki M., Salter-Cid L., Flajnik M.F.;  
 RL J. Immunol. 0:0-0(0).  
 DR EMBL; D87689; BAA19760.1; -.  
 DR MEROPS; T01.010; -.  
 DR INTERPRO; IPR000243; -.  
 DR INTERPRO; IPR001353; -.  
 DR PFAM; PF00227; proteasome; 1.  
 DR PRINTS; PR00141; PROTEASOME.  
 DR PROSITE; PS00854; PROTEASOME.B; 1.  
 KW Proteasome; Hydrolase; Protease.  
 SQ SEQUENCE 230 AA; 24553 MW; 2A1C9B3494473D87 CRC64;

Query Match 4.1%; Score 7; DB 13; Length 230;  
 Best Local Similarity 100.0%; Pred.No.16;  
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 53 TRECLK 59  
 DB 178 TRECLK 184

Search completed: January 31, 2001, 15:07:35  
 Job time: 132 sec

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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:04:19 ; Search time 15.92 seconds  
(without alignments)  
191.752 Million cell updates/sec

Title: US-09-441-654a-1  
Perfect score: 170  
Sequence: 1 ADERSIHDFCLVSKVGRGRC.....ACMLRCFRQENPPLGSK 170

Scoring table: OLIGO  
Gapop 60.0 , Gapext 60.0

Aligned: 174772 seqs, 17957048 residues

Size: 0

Total number of hits satisfying chosen parameters: 174772

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Database : Issued Patents AA: \*  
1: /cgn2.6/ptodata/2/1aa/5A-COMB.pep:\*  
2: /cgn2.6/ptodata/2/1aa/5B-COMB.pep:\*  
3: /cgn2.6/ptodata/2/1aa/6-COMB.pep:\*  
4: /cgn2.6/ptodata/2/1aa/PCRUS-COMB.pep:\*  
5: /cgn2.6/ptodata/2/1aa/backfiles1.pep:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	% Match	Length	ID	Description
1	170	100.0	252	1	US-08-685-660A-7
2	170	100.0	252	1	US-08-974-196-7
3	26	15.3	26	1	US-08-685-660A-3
4	26	15.3	26	2	US-08-974-196-3
5	15	8.8	29	1	US-08-685-660A-2
6	15	8.8	29	2	US-08-974-196-2
7	10	5.9	15	1	US-08-685-660A-1
8	10	5.9	15	2	US-08-974-196-1
9	10	5.9	58	3	US-08-676-124-93
10	10	5.9	58	3	US-08-676-124-103
11	10	5.9	58	3	US-09-414-878-93
12	10	5.9	58	3	US-09-414-878-103
13	10	5.9	58	3	US-09-240-136-93
14	10	5.9	58	3	US-09-240-136-103
15	9	5.3	58	1	US-08-358-160-116
16	9	5.3	58	3	US-08-676-124-111
17	9	5.3	58	3	US-09-414-878-111
18	9	5.3	58	3	US-09-240-136-111
19	9	5.3	59	5	5466783-6
20	9	5.3	65	1	US-08-358-160-92
21	9	5.3	65	5	5466783-12
22	8	4.7	58	1	US-08-463-155A-61
23	8	4.7	58	1	US-08-463-432B-61
24	8	4.7	58	3	US-08-676-124-96
25	8	4.7	58	3	US-09-414-878-96
26	8	4.7	58	3	US-09-240-136-96
27	7	4.1	41	2	US-08-640-847C-41
28	7	4.1	57	1	US-08-358-160-126

29	7	4.1	57	2	US-08-829-876-152	Sequence 152, Appl
30	7	4.1	58	1	US-08-321-658B-4	Sequence 4, Appli
31	7	4.1	58	1	US-08-321-658B-5	Sequence 5, Appli
32	7	4.1	58	1	US-08-463-155A-1	Sequence 1, Appli
33	7	4.1	58	1	US-08-463-155A-2	Sequence 2, Appli
34	7	4.1	58	1	US-08-463-155A-3	Sequence 3, Appli
35	7	4.1	58	1	US-08-463-155A-4	Sequence 4, Appli
36	7	4.1	58	1	US-08-463-155A-45	Sequence 45, Appli
37	7	4.1	58	1	US-08-463-155A-46	Sequence 46, Appli
38	7	4.1	58	1	US-08-463-155A-47	Sequence 47, Appli
39	7	4.1	58	1	US-08-463-155A-48	Sequence 48, Appli
40	7	4.1	58	1	US-08-463-155A-49	Sequence 49, Appli
41	7	4.1	58	1	US-08-463-155A-50	Sequence 50, Appli
42	7	4.1	58	1	US-08-463-155A-51	Sequence 51, Appli
43	7	4.1	58	1	US-08-463-155A-52	Sequence 52, Appli
44	7	4.1	58	1	US-08-463-155A-53	Sequence 53, Appli
45	7	4.1	58	1	US-08-463-155A-54	Sequence 54, Appli

ALIGNMENTS

RESULT 1  
US-08-685-660A-7  
; Sequence 7, Application US/08685660A  
; Patent No. 5731412  
; GENERAL INFORMATION:  
; APPLICANT: SHIMOMURA, Takeshi  
; APPLICANT: KAWAGUCHI, Toshiya  
; APPLICANT: KITAMURA, Naomi  
; TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
; TITLE OF INVENTION: AND METHOD OF PRODUCING THE PROTEIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: SUGHRUE, MIOM, ZINN, MACPEAK & SEAS  
; STREET: 2100 Pennsylvania Avenue, N.W.  
; CITY: Washington  
; STATE: DC  
; COUNTRY: USA  
; ZIP: 20037  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy Disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patentin Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/685,660A  
; FILING DATE: 24-JUL-1996  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: JPA Hei 7-187134  
; FILING DATE: 24-JUL-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Kit, Gordon  
; REGISTRATION NUMBER: 30,764  
; REFERENCE/DOCKET NUMBER: O-42295  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (202) 293-7060  
; TELEFAX: (202) 293-7860  
; INFORMATION FOR SEQ ID NO: 7:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 252 amino acids  
; TYPE: amino acid  
; TOPOLOGY: linear  
; MOLECULE TYPE: peptide  
; US-08-685-660A-7

Query Match 100.0%; Score 170; DB 1; Length 252;  
Best Local Similarity 100.0%; Pred. No. 7, 5e-150; Indels 0; Gaps 0;  
Matches 170; Conservative 0; Mismatches 0;  
QY 1 ADERSIHDFCLVSKVGRGRCASMPRWNVTDGSCQLFVYGGCDGNSNNYLTKECLKK 60  
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DB 28 ADERSIHDCLVSKVGRASMPRWYNTDGSQCLFVYGGCDGNSNNYLTKKECLAK 87  
QY 61 CATVTENATGDLATSRNAADSSVPSAPRQDSEDHSDMFNEEYCTANAVTGPCRASFP 120  
DB 88 CATVTENATGDLATSRNAADSSVPSAPRQDSEDHSDMFNEEYCTANAVTGPCRASFP 147  
QY 121 RWYFDVERNSCNFFIYGGCGNKNYSRSEACMLRCFRQENPPLPLGSK 170  
DB 148 RWYFDVERNSCNFFIYGGCGNKNYSRSEACMLRCFRQENPPLPLGSK 197

RESULT 2  
US-08-974-196-7  
; Sequence 7, Application US/08974196  
; Patent No. 5854396  
; GENERAL INFORMATION:  
; APPLICANT: SHIMOMURA, Takeshi  
; APPLICANT: KAWAGUCHI, Toshiya  
; APPLICANT: KITAMURA, Naomi  
; TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
; TITLE OF INVENTION: AND METHOD OF PRODUCING THE PROTEIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS  
; STREET: 2100 Pennsylvania Avenue, N.W.  
; CITY: Washington  
; STATE: DC  
; COUNTRY: USA  
; ZIP: 20037  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy Disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/974.196  
; FILING DATE: 24-JUL-1996  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: JPA Hei 7-187134  
; FILING DATE: 24-JUL-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: KIT, Gordon  
; REGISTRATION NUMBER: 30,764  
; REFERENCE/DOCKET NUMBER: Q-42295  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (202) 293-7060  
; TELEFAX: (202) 293-7860  
; INFORMATION FOR SEQ ID NO: 7:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 252 amino acids  
; TYPE: amino acid  
; TOPOLOGY: linear  
; MOLECULE TYPE: peptide  
US-08-974-196-7

Query Match 100.0%; Score 170; DB 2; Length 252;  
Best Local Similarity 100.0%; Pred. No. 7.5e-150;  
Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADERSIHDCLVSKVGRASMPRWYNTDGSQCLFVYGGCDGNSNNYLTKKECLAK 60  
DB 28 ADERSIHDCLVSKVGRASMPRWYNTDGSQCLFVYGGCDGNSNNYLTKKECLAK 87  
QY 61 CATVTENATGDLATSRNAADSSVPSAPRQDSEDHSDMFNEEYCTANAVTGPCRASFP 120  
DB 88 CATVTENATGDLATSRNAADSSVPSAPRQDSEDHSDMFNEEYCTANAVTGPCRASFP 147  
QY 121 RWYFDVERNSCNFFIYGGCGNKNYSRSEACMLRCFRQENPPLPLGSK 170  
DB 148 RWYFDVERNSCNFFIYGGCGNKNYSRSEACMLRCFRQENPPLPLGSK 197

US-08-685-660A-3  
; Sequence 3, Application US/08685660A  
; Patent No. 5731412  
; GENERAL INFORMATION:  
; APPLICANT: SHIMOMURA, Takeshi  
; APPLICANT: KAWAGUCHI, Toshiya  
; APPLICANT: KITAMURA, Naomi  
; TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
; TITLE OF INVENTION: AND METHOD OF PRODUCING THE PROTEIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS  
; STREET: 2100 Pennsylvania Avenue, N.W.  
; CITY: Washington  
; STATE: DC  
; COUNTRY: USA  
; ZIP: 20037  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy Disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/685,660A  
; FILING DATE: 24-JUL-1996  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: JPA Hei 7-187134  
; FILING DATE: 24-JUL-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: KIT, Gordon  
; REGISTRATION NUMBER: 30,764  
; REFERENCE/DOCKET NUMBER: Q-42295  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (202) 293-7060  
; TELEFAX: (202) 293-7860  
; INFORMATION FOR SEQ ID NO: 3:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 26 amino acids  
; TYPE: amino acid  
; TOPOLOGY: linear  
; MOLECULE TYPE: peptide  
; FRAGMENT TYPE: internal fragment  
; ORIGINAL SOURCE:  
; ORGANISM: Homo sapiens  
; STRAIN: MKN45  
US-08-685-660A-3

Query Match 15.3%; Score 26; DB 1; Length 26;  
Best Local Similarity 100.0%; Pred. No. 1.4e-17;  
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 62 ATVTENATGDLATSRNAADSSVPSAP 87  
DB 1 ATVTENATGDLATSRNAADSSVPSAP 26

RESULT 4  
US-08-974-196-3  
; Sequence 3, Application US/08974196  
; Patent No. 5854396  
; GENERAL INFORMATION:  
; APPLICANT: SHIMOMURA, Takeshi  
; APPLICANT: KAWAGUCHI, Toshiya  
; APPLICANT: KITAMURA, Naomi  
; TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
; TITLE OF INVENTION: AND METHOD OF PRODUCING THE PROTEIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS



STREET: 2100 Pennsylvania Avenue, N.W.  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20037  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy Disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/974,196  
FILING DATE: 24-JUL-1996  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/685,660  
FILING DATE: 24-JUL-1996  
APPLICATION NUMBER: JPA Hei 7-187134  
FILING DATE: 24-JUL-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: KIT, Gordon  
REGISTRATION NUMBER: 30,764  
REFERENCE/DOCKET NUMBER: Q-42295  
TELEPHONE: (202) 293-7060  
TELEFAX: (202) 293-7860  
INFORMATION FOR SEQ ID NO: 3:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 26 amino acids  
TYPE: amino acid  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
FRAGMENT TYPE: internal fragment  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
STRAIN: MKN45  
US-08-974-196-3

Query Match 15.3%; Score 26; DB 2; Length 26;  
Best Local Similarity 100.0%; Pred. No. 1.4e-17;  
Matches 26; Conservative 0; Mismatches 0; Indels 0;

QY 62 ATVTENATGDLATSRNADSSVSPAP 87  
|||||  
Db 1 ATVTENATGDLATSRNADSSVSPAP 26  
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SULT 5  
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Sequence 2, Application US/08685660A  
Patent No. 5731412  
GENERAL INFORMATION:  
APPLICANT: SHIMOMURA, Takeshi  
APPLICANT: KAWAGUCHI, Toshiya  
APPLICANT: KITAMURA, Naomi  
TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
NUMBER OF SEQUENCES: 7  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS  
STREET: 2100 Pennsylvania Avenue, N.W.  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20037  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy Disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/685,660A  
FILING DATE: 24-JUL-1996

PRIOR APPLICATION DATA:  
APPLICATION NUMBER: JPA Hei 7-187134  
FILING DATE: 24-JUL-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: KIT, Gordon  
REGISTRATION NUMBER: 30,764  
REFERENCE/DOCKET NUMBER: Q-42295  
TELEPHONE: (202) 293-7060  
TELEFAX: (202) 293-7860  
INFORMATION FOR SEQ ID NO: 2:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 29 amino acids  
TYPE: amino acid  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
FRAGMENT TYPE: internal fragment  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
STRAIN: MKN45  
US-08-685-660A-2

Query Match 8.8%; Score 15; DB 1; Length 29;  
Best Local Similarity 100.0%; Pred. No. 2.2e-07;  
Matches 15; Conservative 0; Mismatches 0; Indels 0;

QY 21 RASMPRWYNNVTDGS 35  
|||||  
Db 7 RASMPRWYNNVTDGS 21  
|||||

RESULT 6  
US-08-974-196-2  
Sequence 2, Application US/08974196  
Patent No. 5854396  
GENERAL INFORMATION:  
APPLICANT: SHIMOMURA, Takeshi  
APPLICANT: KAWAGUCHI, Toshiya  
APPLICANT: KITAMURA, Naomi  
TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
NUMBER OF SEQUENCES: 7  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS  
STREET: 2100 Pennsylvania Avenue, N.W.  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20037  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy Disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/974,196  
FILING DATE:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/685,660  
FILING DATE: 24-JUL-1996  
APPLICATION NUMBER: JPA Hei 7-187134  
FILING DATE: 24-JUL-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: KIT, Gordon  
REGISTRATION NUMBER: 30,764  
REFERENCE/DOCKET NUMBER: Q-42295  
TELEPHONE: (202) 293-7060  
TELEFAX: (202) 293-7860  
INFORMATION FOR SEQ ID NO: 2:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 29 amino acids

us-09-441-654a-1.oli.ra1

wed Jan 31 15:14:32 2001

;; TYPE: amino acid  
;; TOPOLOGY: linear  
;; MOLECULE TYPE: peptide  
;; FRAGMENT TYPE: internal fragment  
;; ORIGINAL SOURCE:  
;; ORGANISM: Homo sapiens  
;; STRAIN: MKN45  
US-08-974-196-2

Query Match 8.8%; Score 15; DB 2; Length 29;  
Best Local Similarity 100.0%; Pred. No. 2.2e-07;  
Matches 15; Conservative 0; Mismatches 0; Indels 0;

QY 21 RASMPRWYNTDGS 35  
DB 7 RASMPRWYNTDGS 21

RESULT 7  
US-08-685-660A-1  
Sequence 1, Application US/08685660A  
Patent No. 5731412

;; GENERAL INFORMATION:  
;; APPLICANT: SHIMOMURA, Takeshi  
;; APPLICANT: KAWAGUCHI, Toshiya  
;; APPLICANT: KITAMURA, Naomi  
;; TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
;; TITLE OF INVENTION: AND METHOD OF PRODUCING THE PROTEIN  
;; NUMBER OF SEQUENCES: 7  
;; CORRESPONDENCE ADDRESS:  
;; ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS  
;; STREET: 2100 Pennsylvania Avenue, N.W.  
;; CITY: Washington  
;; STATE: DC  
;; COUNTRY: USA  
;; ZIP: 20037

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy Disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/685,660A  
FILING DATE: 24-JUL-1996  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: JPA Hei 7-187134  
FILING DATE: 24-JUL-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: KIT, Gordon

REGISTRATION NUMBER: 30,764  
REFERENCE/DOCKET NUMBER: Q-42295  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (202) 293-7060  
TELEFAX: (202) 293-7860  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 15 amino acids  
TYPE: amino acid  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
FRAGMENT TYPE: N-terminal fragment  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
STRAIN: MKN45  
US-08-685-660A-1

Query Match 5.9%; Score 10; DB 1; Length 15;  
Best Local Similarity 100.0%; Pred. No. 0.0053;  
Matches 10; Conservative 0; Mismatches 0; Indels 0;

QY 1 ADERSIHDF 10

DB 1 ADERSIHDF 10

RESULT 8

US-08-974-196-1  
Sequence 1, Application US/08974196  
Patent No. 5854396  
GENERAL INFORMATION:  
APPLICANT: SHIMOMURA, Takeshi  
APPLICANT: KAWAGUCHI, Toshiya  
APPLICANT: KITAMURA, Naomi  
TITLE OF INVENTION: NOVEL PROTEIN, DNA CODING FOR SAME  
TITLE OF INVENTION: AND METHOD OF PRODUCING THE PROTEIN  
NUMBER OF SEQUENCES: 7  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS  
STREET: 2100 Pennsylvania Avenue, N.W.  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20037

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy Disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/974,196  
FILING DATE:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/685,660  
FILING DATE: 24-JUL-1996  
APPLICATION NUMBER: JPA Hei 7-187134  
FILING DATE: 24-JUL-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: KIT, Gordon

REGISTRATION NUMBER: 30,764  
REFERENCE/DOCKET NUMBER: Q-42295  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (202) 293-7060  
TELEFAX: (202) 293-7860  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 15 amino acids  
TYPE: amino acid  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
FRAGMENT TYPE: N-terminal fragment  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
STRAIN: MKN45  
US-08-974-196-1

Query Match 5.9%; Score 10; DB 2; Length 15;  
Best Local Similarity 100.0%; Pred. No. 0.0053;  
Matches 10; Conservative 0; Mismatches 0; Indels 0;

QY 1 ADERSIHDF 10  
DB 1 ADERSIHDF 10

RESULT 9

US-08-676-124-93  
Sequence 93, Application US/08676124  
Patent No. 6010880  
GENERAL INFORMATION:  
APPLICANT: MARKLAND, William  
APPLICANT: LADNER, Robert Charles  
TITLE OF INVENTION: INHIBITORS OF HUMAN PLASMIN DERIVED  
TITLE OF INVENTION: FROM FROM THE KUNITZ DOMAINS

NUMBER OF SEQUENCES: 137  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Browdy and Neimark  
STREET: 419 Seventh Street N.W., Ste. 300  
CITY: Washington  
STATE: D.C.  
COUNTRY: United States of America  
ZIP: 20004  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/676,124  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/00298  
FILING DATE: 11-JAN-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/179,658  
FILING DATE: 11-JAN-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/208,265  
FILING DATE: 10-MAR-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: COOPER, IVER P.  
REGISTRATION NUMBER: 28,005  
REFERENCE/DOCKET NUMBER: MARKLAND=3B  
TELEPHONE: 202-628-5197  
TELEFAX: 202-737-3528  
INFORMATION FOR SEQ ID NO: 93:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 58 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
US-08-676-124-93

Query Match 5.9%; Score 10; DB 3; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.016;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

113 GPCRASFPRW 122  
|||||

12 GPCRASFPRW 21

RESULT 10  
US-08-676-124-103  
Sequence 103, Application US/08676124  
Patent No. 6010880  
GENERAL INFORMATION:  
APPLICANT: MARKLAND, William  
ADDRESSEE: LADNER, Robert Charles  
TITLE OF INVENTION: INHIBITORS OF HUMAN PLASMIN DERIVED  
FROM FROM THE KUNITZ DOMAINS  
NUMBER OF SEQUENCES: 137  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Browdy and Neimark  
STREET: 419 Seventh Street N.W., Ste. 300  
CITY: Washington  
STATE: D.C.  
COUNTRY: United States of America  
ZIP: 20004  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/676,124  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/00298  
FILING DATE: 11-JAN-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/179,658  
FILING DATE: 11-JAN-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/208,265  
FILING DATE: 10-MAR-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: COOPER, IVER P.  
REGISTRATION NUMBER: 28,005  
REFERENCE/DOCKET NUMBER: MARKLAND=3B  
TELEPHONE: 202-628-5197  
TELEFAX: 202-737-3528  
INFORMATION FOR SEQ ID NO: 103:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 58 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
US-08-676-124-103

Query Match 5.9%; Score 10; DB 3; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.016;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

112 TGPCRASFPR 121  
|||||

11 TGPCRASFPR 20

RESULT 11  
US-09-414-878-93  
Sequence 93, Application US/09414878  
Patent No. 6071723  
GENERAL INFORMATION:  
APPLICANT: DYAX CORP  
ADDRESSEE: Yankwich & Associates  
TITLE OF INVENTION: Inhibitors of Human Plasmin Derived  
FROM THE KUNITZ DOMAINS  
NUMBER OF SEQUENCES: 139  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Yankwich & Associates  
STREET: 130 Bishop Allen Drive  
CITY: Cambridge  
STATE: Massachusetts  
COUNTRY: USA  
ZIP: 02139  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5-inch diskette  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: Microsoft Windows 98  
SOFTWARE: Microsoft Word 97 SR-1  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/414,878  
FILING DATE: (concurrently herewith)  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 09/240,136  
FILING DATE: 29-JAN-1999  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/676,124  
FILING DATE: 07-JAN-1997

; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: PCT/US95/00298  
; FILING DATE: 11-JAN-1995  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/208,265  
; FILING DATE: 10-MAR-1994  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/179,685  
; FILING DATE: 11-JAN-1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: YANKWICH, Leon R  
; REGISTRATION NUMBER: 30,237  
; REFERENCE/DOCKET NUMBER: DYX-007.2P US-2  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 617-491-8801  
; TELEFAX: 617-491-8801  
; INFORMATION FOR SEQ ID NO: 93:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 58 amino acids  
; TYPE: amino acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: protein  
; US-09-414-878-93

Query Match 5.9%; Score 10; DB 3; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.016;  
Matches 10; Conservative 0; Mismatches 0; Indels 0;

QY 113 GPCRSPRPW 122  
Db 12 GPCRSPRPW 21  
|||||

RESULT 12  
US-09-414-878-103  
; Sequence 103, Application US/09414878  
; Patent No. 6071723  
; GENERAL INFORMATION:  
; APPLICANT: DYAX CORP  
; APPLICANT: MARKLAND, William  
; APPLICANT: LADNER, Robert C  
; TITLE OF INVENTION: Inhibitors of Human Plamin Derived  
; TITLE OF INVENTION: From The Kunitz Domains  
; NUMBER OF SEQUENCES: 139  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Yankwich & Associates  
; STREET: 130 Bishop Allen Drive  
; CITY: Cambridge  
; STATE: Massachusetts  
; COUNTRY: USA  
; ZIP: 02139  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: 3.5-inch diskette  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: Microsoft Windows 98  
; SOFTWARE: Microsoft Word 97 SR-1  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/414,878  
; FILING DATE: (concurrently herewith)  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 09/240,136  
; FILING DATE: 29-JAN-1999  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/676,124  
; FILING DATE: 07-JAN-1997  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: PCT/US95/00298  
; FILING DATE: 11-JAN-1995

; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/208,265  
; FILING DATE: 10-MAR-1994  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/179,685  
; FILING DATE: 11-JAN-1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: YANKWICH, Leon R  
; REGISTRATION NUMBER: 30,237  
; REFERENCE/DOCKET NUMBER: DYX-007.2P US-2  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 617-491-4343  
; TELEFAX: 617-491-8801  
; INFORMATION FOR SEQ ID NO: 103:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 58 amino acids  
; TYPE: amino acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: protein  
; US-09-414-878-103

Query Match 5.9%; Score 10; DB 3; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.016;  
Matches 10; Conservative 0; Mismatches 0; Indels 0;

QY 112 TGPCRASFPR 121  
Db 11 TGPCRASFPR 20  
|||||

RESULT 13  
US-09-240-136-93  
; Sequence 93, Application US/09240136  
; Patent No. 6103499  
; GENERAL INFORMATION:  
; APPLICANT: DYAX CORP  
; APPLICANT: MARKLAND, William  
; APPLICANT: LADNER, Robert C  
; TITLE OF INVENTION: Inhibitors of Human Plamin Derived  
; TITLE OF INVENTION: From The Kunitz Domains  
; NUMBER OF SEQUENCES: 139  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Yankwich & Associates  
; STREET: 130 Bishop Allen Drive  
; CITY: Cambridge  
; STATE: Massachusetts  
; COUNTRY: USA  
; ZIP: 02139  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: 3.5-inch diskette  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: Microsoft Windows 98  
; SOFTWARE: Microsoft Word 97 SR-1  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/240,136  
; FILING DATE: (concurrently herewith)  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/676,124  
; FILING DATE: 07-JAN-1997  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: PCT/US95/00298  
; FILING DATE: 11-JAN-1995  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/208,265  
; FILING DATE: 10-MAR-1994  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/179,685  
; FILING DATE: 11-JAN-1994

ATTORNEY/AGENT INFORMATION:  
NAME: YANKWICH, Leon R  
REGISTRATION NUMBER: 30,237  
NAME: ZWICKER, Kenneth P  
REGISTRATION NUMBER: 43,310  
REFERENCE/DOCKET NUMBER: DYX-007.2P US-1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 617-491-4343  
TELEFAX: 617-491-8801  
INFORMATION FOR SEQ ID NO: 93:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 58 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
US-09-240-136-93

Query Match 5.9%; Score 10; DB 3; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.016;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 113 GPCRASPPRW 122  
Db 12 GPCRASPPRW 21

RESULT 14  
US-09-240-136-103  
Sequence 103, Application US/09240136  
Patent No. 6103499  
GENERAL INFORMATION:  
APPLICANT: DYAX CORP  
APPLICANT: LADNER, Robert C  
APPLICANT: MARKLAND, William  
TITLE OF INVENTION: Inhibitors of Human Plamin Derived  
TITLE OF INVENTION: From The Kunitz Domains  
NUMBER OF SEQUENCES: 139  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Yankwich & Associates  
STREET: 130 Bishop Allen Drive  
CITY: Cambridge  
STATE: Massachusetts  
COUNTRY: USA  
ZIP: 02139  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5-inch diskette  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: Microsoft Windows 98  
SOFTWARE: Microsoft Word 97 SR-1  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/240,136  
FILING DATE: (concurrently herewith)  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/676,124  
FILING DATE: 07-JAN-1997  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/00298  
FILING DATE: 11-JAN-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/208,265  
FILING DATE: 10-MAR-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/179,695  
FILING DATE: 11-JAN-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: YANKWICH, Leon R  
REGISTRATION NUMBER: 30,237  
NAME: ZWICKER, Kenneth P  
REGISTRATION NUMBER: 43,310  
REFERENCE/DOCKET NUMBER: DYX-007.2P US-1

TELECOMMUNICATION INFORMATION:  
TELEPHONE: 617-491-4343  
TELEFAX: 617-491-8801  
INFORMATION FOR SEQ ID NO: 103:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 58 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
US-09-240-136-103  
Query Match 5.9%; Score 10; DB 3; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.016;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 112 TGPCRASFP 121  
Db 11 TGPCRASFP 20  
RESULT 15  
US-08-358-160-116  
Sequence 116, Application US/08358160  
Patent No. 5663143  
GENERAL INFORMATION:  
APPLICANT: LEY, Arthur C.  
APPLICANT: LADNER, Robert C.  
APPLICANT: GUTERMAN, Sonia K.  
APPLICANT: ROBERTS, Bruce L.  
APPLICANT: MARKLAND, William  
APPLICANT: KENT, Rachel B.  
TITLE OF INVENTION: ENGINEERED HUMAN-DERIVED KUNITZ  
TITLE OF INVENTION: DOMAINS THAT INHIBIT HUMAN NEUTROPHIL ELASTASE  
NUMBER OF SEQUENCES: 234  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: BROWDY AND NEIMARK  
STREET: 419 Seventh Street, N.W. Suite 300  
CITY: Washington  
STATE: District of Columbia  
COUNTRY: USA  
ZIP: 20004  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/358,160  
FILING DATE: 16-DEC-1994  
CLASSIFICATION: 514  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/133,031  
FILING DATE: 13-OCT-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/009,319  
FILING DATE: 26-JAN-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/664,989  
FILING DATE: 01-MAR-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/487,063  
FILING DATE: 02-MAR-1990  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/240,160  
FILING DATE: 02-SEP-1988  
ATTORNEY/AGENT INFORMATION:  
NAME: Cooper, Iver P.  
REGISTRATION NUMBER: 28,005  
REFERENCE/DOCKET NUMBER: LEY-1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-628-5197

; TELEFAX: 202-737-3528  
; TELEX: 248633  
; INFORMATION FOR SEQ ID NO: 116:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 58 amino acids  
; TYPE: amino acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: protein  
; US-08-358-160-116

Query Match 5.3%; Score 9; DB 1; Length 58;  
Best Local Similarity 100.0%; Pred. No. 0.13;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 FIYGGCRGN 142  
| | | | |  
Db 35 FIYGGCRGN 43

Search completed: January 31, 2001, 15:06:02  
Job time: 103 sec

GenCore version 4.5  
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OM protein - protein search, using sw model

Run on: January 31, 2001, 15:04:16 ; Search time 16.78 seconds  
(without alignments)  
346.421 Million cell updates/sec

Title: US-09-441-654A-1

Perfect score: 170

Sequence: 1 ADERSIHDFCLVSKVGRG.....ACMLRCFRQENPPLPLGSK 170

Scoring table: OLIGO

Gapop 60.0 , Gapext 60.0

Searched: 268485 seqs, 34193795 residues

d size : 0

Total number of hits satisfying chosen parameters: 268485

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

- A\_Geneseq\_36:\*
- 1: /SIDS1/gcgdata/geneseq/geneseq/AA1980.DAT:\*
  - 2: /SIDS1/gcgdata/geneseq/geneseq/AA1981.DAT:\*
  - 3: /SIDS1/gcgdata/geneseq/geneseq/AA1982.DAT:\*
  - 4: /SIDS1/gcgdata/geneseq/geneseq/AA1983.DAT:\*
  - 5: /SIDS1/gcgdata/geneseq/geneseq/AA1984.DAT:\*
  - 6: /SIDS1/gcgdata/geneseq/geneseq/AA1985.DAT:\*
  - 7: /SIDS1/gcgdata/geneseq/geneseq/AA1986.DAT:\*
  - 8: /SIDS1/gcgdata/geneseq/geneseq/AA1987.DAT:\*
  - 9: /SIDS1/gcgdata/geneseq/geneseq/AA1988.DAT:\*
  - 10: /SIDS1/gcgdata/geneseq/geneseq/AA1989.DAT:\*
  - 11: /SIDS1/gcgdata/geneseq/geneseq/AA1990.DAT:\*
  - 12: /SIDS1/gcgdata/geneseq/geneseq/AA1991.DAT:\*
  - 13: /SIDS1/gcgdata/geneseq/geneseq/AA1992.DAT:\*
  - 14: /SIDS1/gcgdata/geneseq/geneseq/AA1993.DAT:\*
  - 15: /SIDS1/gcgdata/geneseq/geneseq/AA1994.DAT:\*
  - 16: /SIDS1/gcgdata/geneseq/geneseq/AA1995.DAT:\*
  - 17: /SIDS1/gcgdata/geneseq/geneseq/AA1996.DAT:\*
  - 18: /SIDS1/gcgdata/geneseq/geneseq/AA1997.DAT:\*
  - 19: /SIDS1/gcgdata/geneseq/geneseq/AA1998.DAT:\*
  - 20: /SIDS1/gcgdata/geneseq/geneseq/AA1999.DAT:\*
  - 21: /SIDS1/gcgdata/geneseq/geneseq/AA2000.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	170	100.0	170	W30041	Human placental bi
2	170	100.0	179	W30053	Human placental bi
3	170	100.0	197	W30043	Human placental bi
4	170	100.0	213	W30042	Human placental bi
5	170	100.0	225	W30046	Human placental bi
6	170	100.0	235	W30060	Human placental bi
7	170	100.0	240	W30045	Human placental bi
8	170	100.0	248	W30044	Human placental bi
9	170	100.0	252	W30040	Human placental bi
10	170	100.0	252	W13665	Hepatocyte growth
11	170	100.0	252	W70286	Human tissue facto
12	153	90.0	153	W30051	Human placental bi

13	146	85.9	146	18	W30052	Human placental bi
14	92	54.1	92	18	W30054	Human placental bi
15	84	49.4	169	18	W30063	EST R74593 protein
16	72	42.4	130	18	W30062	EST R35464 protein
17	58	34.1	58	18	W30047	Human placental bi
18	58	34.1	58	18	W30049	Human placental bi
19	51	30.0	51	18	W30048	Human placental bi
20	51	30.0	51	18	W30050	Human placental bi
21	38	22.4	170	18	W30061	Human consensus bi
22	26	15.3	26	18	W13664	Hepatocyte growth
23	15	8.8	29	18	W13663	Hepatocyte growth
24	11	6.5	302	14	R35001	LACI. Rattus rat
25	11	6.5	302	17	R88513	Lipoprotein-associ
26	10	5.9	12	16	R78576	LACI K1 derivative
27	10	5.9	12	16	R78586	LACI K1 derivative
28	10	5.9	15	18	W13662	Hepatocyte growth
29	9	5.3	12	16	R78594	LACI K1 derivative
30	8	4.7	12	16	R78579	LACI K1 derivative
31	8	4.7	58	18	W07766	Non-native Kunitz-
32	8	4.7	58	19	W64138	Human Kunitz-type
33	8	4.7	219	21	Y54090	Enzyme EFSE involv
34	8	4.7	219	21	Y43792	Amino acid sequenc
35	7	4.1	12	16	R78578	LACI K1 derivative
36	7	4.1	56	14	R39677	C-terminal Kunitz-
37	7	4.1	57	11	R08293	Example of Alzheim
38	7	4.1	57	19	W47434	Aprotinin variant
39	7	4.1	57	19	W47435	Aprotinin variant
40	7	4.1	57	19	W47436	Aprotinin variant
41	7	4.1	57	21	Y68103	Kunitz protease in
42	7	4.1	58	14	R39673	C-terminal Kunitz-
43	7	4.1	58	14	R39799	Kunitz-type protea
44	7	4.1	58	14	R39800	Kunitz-type protea
45	7	4.1	58	16	R78556	Human III-Kudow 2

ALIGNMENTS

RESULT 1  
W30041  
ID W30041 standard; Protein; 170 AA.  
XX W30041;

DT 20-APR-1998 (first entry)

DE Human placental bikunin.

Human; placental bikunin; inhibition; trypsin; kallikrein;  
plasmin; factor XIIa; treatment; prevention; oedema;  
inflammation; infection; granulomatosis; multiple sclerosis;  
ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
blood coagulation disease; polytrauma; stroke; haemorrhage;  
gastric cancer; cervical cancer; metastasis; blood loss.

OS Homo sapiens.

PN WO9733996-A2.

PD 18-SEP-1997.

PF 10-MAR-1997; 97WO-US03894.

PR 04-OCT-1996; 96US-0725251.

PR 11-MAR-1996; 96US-0013106.

PR 14-JUN-1996; 96US-0019793.

PA (FARB ) BAYER CORP.

PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP:

DR WPI; 1997-470876/43.

XX

PT New human placental bikunin - used to inhibit kallikrein, trypsin  
 PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 PT perioperative blood loss

XX Claim 1; Page 65; 110pp; English.

XX The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.

XX Sequence 170 AA;

Query Match 100.0%; Score 170; DB 18; Length 170;  
 . Best Local Similarity 100.0%; Pred. No. 2.4e-156;

. Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADERSIHDFCLVSKVGRCRASPRWYNYVDGSCOLFVYGGDGNNSNYLTKEECLKK 60  
 QD 1 adersihdfclvskvgrcrasprwvnyvtdgscqlfvyggcdgnsnyltkeecikl 60  
 QY 61 CATVTENATGDLATSRNAADSSVPSAPRRQDSEHSDMFNEEYCTANAVTGPCRASFP 120  
 DB 61 catvtenatgdlatsrnaadssvpsaprrqdsedhssdmfneyeyctananavtgcra 120  
 QY 121 RWFYDVERNSNNFIYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK 170  
 DB 121 rwyfdvernsnnfiyggcrgnknsyrseacmlrcfrqenpplplgsk 170

RESULT 2  
 W30053 W30053 standard; Protein; 179 AA.

AC W30053;

XX 20-APR-1998 (first entry)

XX Human placental bikunin.

XX Human; placental bikunin; inhibition; trypsin; kallikrein;  
 KW plasmin; factor XIIa; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.

XX Homo sapiens.

XX WO9733996-A2.

XX 18-SEP-1997.

XX 10-MAR-1997; 97WO-US03894.

XX 04-OCT-1996; 96US-0725251.

PR 11-MAR-1996; 96US-0013106.

PR 14-JUN-1996; 96US-0019793.

XX (FARB ) BAYER CORP.

PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;

XX WPI; 1997-470876/43.

XX New human placental bikunin - used to inhibit kallikrein, trypsin  
 PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 PT perioperative blood loss

XX Claim 1; Page 67; 110pp; English.

XX The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.

XX Sequence 179 AA;

Query Match 100.0%; Score 170; DB 18; Length 179;  
 Best Local Similarity 100.0%; Pred. No. 2.5e-156;

. Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADERSIHDFCLVSKVGRCRASPRWYNYVDGSCOLFVYGGDGNNSNYLTKEECLKK 60  
 DB 1 adersihdfclvskvgrcrasprwvnyvtdgscqlfvyggcdgnsnyltkeecikl 60  
 QY 61 CATVTENATGDLATSRNAADSSVPSAPRRQDSEHSDMFNEEYCTANAVTGPCRASFP 120  
 DB 61 catvtenatgdlatsrnaadssvpsaprrqdsedhssdmfneyeyctananavtgcra 120  
 QY 121 RWFYDVERNSNNFIYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK 170  
 DB 121 rwyfdvernsnnfiyggcrgnknsyrseacmlrcfrqenpplplgsk 170

RESULT 3

W30043 W30043 standard; Protein; 197 AA.

XX W30043;

XX 20-APR-1998 (first entry)

XX Human placental bikunin.

XX Human; placental bikunin; inhibition; trypsin; kallikrein;  
 KW plasmin; factor XIIa; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.

XX Homo sapiens.



W09733996-A2.  
 18-SEP-1997.  
 10-MAR-1997; 97WO-US03894.  
 04-OCT-1996; 96US-0725251.  
 11-MAR-1996; 96US-0013106.  
 14-JUN-1996; 96US-0019793.  
 (FARB ) BAYER CORP.

Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 WPI; 1997-470876/43.

New human placental bikunin - used to inhibit kallikrein, trypsin  
 etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 perioperative blood loss

Claim 1; Page 65; 110pp; English.

The present sequence is a human placental bikunin, which  
 inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
 Bikunin can be used to treat or prevent brain and spinal cord  
 oedema, inflammation, infection or granulomatosis, multiple  
 sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 fibrosis, blood coagulation diseases, polytrauma, stroke,  
 cerebral or subarachnoid haemorrhage and gastric or cervical  
 cancer and prevent metastasis. It is particularly useful for  
 reducing blood loss during surgery, and can also be used to treat  
 other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 influenza and similar viral infections, acute pancreatitis and  
 gout, and prevent pre-term labour. It has similar properties to  
 aprotinin, but is less highly charged so should be less  
 immunogenic and less likely to damage the kidneys. Manipulation  
 of the bikunin sequence may allow the inhibitory profile to be  
 altered. It also reduces or eliminates the need for whole donor  
 blood or blood products during surgery, thereby reducing the risk  
 of infection and other adverse side effects, as well as reducing  
 the cost of surgery.

Sequence 197 AA;

Query Match 100.0%; Score 170; DB 18; Length 197;  
 Best Local Similarity 100.0%; Pred. No. 2.7e-156;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 ADERSIHDFCLVSKVVGRCRASPMPRWYNTDGSQOLFVYGGCDGNSNNYLTKEECLKK 60  
 19 adersihdfclvskvvgrcrasmprrwyntdgsqolfvyygdcgnsnnyltkeecclk 78  
 61 CATVTENATGDLATSRNAADSSVPSAPPRODSEDHSDMFNEYEYCTANAVTGPCRASFP 120  
 79 catvtenatgdlatsrnaadssvpsaprrqdsdhsdmfneyeyctanavtgpccrasfp 138  
 121 RWYFDVERNSCNFFYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK 170  
 139 rwyfdvernsccnnfiyggcrgnknysrseacmlrcfrqenpplplgsk 188

RESULT 4  
 W30042  
 ID W30042 standard; Protein; 213 AA.  
 AC W30042;  
 XX 20-APR-1998 (first entry)  
 XX Human placental bikunin.  
 XX Human; placental bikunin; inhibition; trypsin; kallikrein;

KW plasmin; factor XIIa; treatment; prevention; oedema;  
 KW inflammation; infection; granulomatosis; multiple sclerosis;  
 KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
 KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
 KW gastric cancer; cervical cancer; metastasis; blood loss.  
 XX Homo sapiens.  
 OS W09733996-A2.  
 PN 18-SEP-1997.  
 XX 10-MAR-1997;  
 PD 04-OCT-1996;  
 PF 11-MAR-1996;  
 XX 14-JUN-1996;  
 PA (FARB ) BAYER CORP.  
 XX Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 XX WPI; 1997-470876/43.  
 XX New human placental bikunin - used to inhibit kallikrein, trypsin  
 etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 perioperative blood loss  
 PS Claim 1; Page 65; 110pp; English.  
 XX The present sequence is a human placental bikunin, which  
 CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
 CC Bikunin can be used to treat or prevent brain and spinal cord  
 CC oedema, inflammation, infection or granulomatosis, multiple  
 CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 CC fibrosis, blood coagulation diseases, polytrauma, stroke,  
 CC cerebral or subarachnoid haemorrhage and gastric or cervical  
 CC cancer and prevent metastasis. It is particularly useful for  
 CC reducing blood loss during surgery, and can also be used to treat  
 CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 CC influenza and similar viral infections, acute pancreatitis and  
 CC gout, and prevent pre-term labour. It has similar properties to  
 CC aprotinin, but is less highly charged so should be less  
 CC immunogenic and less likely to damage the kidneys. Manipulation  
 CC of the bikunin sequence may allow the inhibitory profile to be  
 CC altered. It also reduces or eliminates the need for whole donor  
 CC blood or blood products during surgery, thereby reducing the risk  
 CC of infection and other adverse side effects, as well as reducing  
 CC the cost of surgery.  
 XX Sequence 213 AA;

Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
 WPI; 1997-470876/43.

New human placental bikunin - used to inhibit kallikrein, trypsin  
 etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
 perioperative blood loss

Claim 1; Page 65; 110pp; English.

The present sequence is a human placental bikunin, which  
 inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
 Bikunin can be used to treat or prevent brain and spinal cord  
 oedema, inflammation, infection or granulomatosis, multiple  
 sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
 fibrosis, blood coagulation diseases, polytrauma, stroke,  
 cerebral or subarachnoid haemorrhage and gastric or cervical  
 cancer and prevent metastasis. It is particularly useful for  
 reducing blood loss during surgery, and can also be used to treat  
 other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
 influenza and similar viral infections, acute pancreatitis and  
 gout, and prevent pre-term labour. It has similar properties to  
 aprotinin, but is less highly charged so should be less  
 immunogenic and less likely to damage the kidneys. Manipulation  
 of the bikunin sequence may allow the inhibitory profile to be  
 altered. It also reduces or eliminates the need for whole donor  
 blood or blood products during surgery, thereby reducing the risk  
 of infection and other adverse side effects, as well as reducing  
 the cost of surgery.

Sequence 213 AA;

Query Match 100.0%; Score 170; DB 18; Length 213;  
 Best Local Similarity 100.0%; Pred. No. 2.9e-156;  
 Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADERSIHDFCLVSKVVGRCRASPMPRWYNTDGSQOLFVYGGCDGNSNNYLTKEECLKK 60  
 Db 1 adersihdfclvskvvgrcrasmprrwyntdgsqolfvyygdcgnsnnyltkeecclk 60  
 QY 61 CATVTENATGDLATSRNAADSSVPSAPPRODSEDHSDMFNEYEYCTANAVTGPCRASFP 120  
 Db 61 catvtenatgdlatsrnaadssvpsaprrqdsdhsdmfneyeyctanavtgpccrasfp 120  
 QY 121 RWYFDVERNSCNFFYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK 170  
 Db 121 rwyfdvernsccnnfiyggcrgnknysrseacmlrcfrqenpplplgsk 170

RESULT 5  
 W30046  
 ID W30046 standard; Protein; 225 AA.



CC blood or blood products during surgery, thereby reducing the risk  
CC of infection and other adverse side effects, as well as reducing  
CC the cost of surgery.

XX Sequence 235 AA;

Query Match 100.0%; Score 170; DB 18; Length 235;  
Best Local Similarity 100.0%; Pred. No. 3.1e-156;  
Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADERSIHDFCLVSKVVGCRASMPRWYNTDSCOLFVYGGDGNNSNYLTKEECLKK 60  
Db 19 adersihdfclvskvvgcrasmprrwvntdgsclfvvgcdgnsnyltkeecclkk 78  
QY 61 CATVTENATGDLATSRNAADSSVPSAPRRQDSEHSDMFNEEYCTANAVTGPCRASFP 120  
Db 79 catvtenatgdlatsrnaadssvpsaprrqdsedhssdmfneeyctanavtgcrafp 138  
121 RWYFDVERNSCNFIYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK 170  
139 rwyfdvernsnfnfiyggcrngknsyrseacmlrcfrqenpplplgsk 188

RESULT 7

ID W30045 standard; Protein: 240 AA.

AC W30045;

DT 20-APR-1998 (first entry)

DE Human placental bikunin.

KW Human; placental bikunin; inhibition; trypsin; kallikrein;  
KW plasmin; factor XIIa; treatment; prevention; oedema;  
KW inflammation; infection; granulomatosis; multiple sclerosis;  
KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
KW gastric cancer; cervical cancer; metastasis; blood loss.

OS Homo sapiens.

PN W09733996-A2.

PD 18-SEP-1997.

PF 10-MAR-1997; 97WO-US03894.

PR 04-OCT-1996; 96US-0725251.

PR 11-MAR-1996; 96US-0013106.

PR 14-JUN-1996; 96US-0019793.

PA (FARB ) BAYER CORP.

XX Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;

XX WPI; 1997-470876/43.

XX N-PSDB; T90734.

XX New human placental bikunin - used to inhibit kallikrein, trypsin

XX etc. in treatment of oedema, multiple sclerosis, fibrosis, or

XX perioperative blood loss

XX Claim 1; Page 66; 110pp; English.

XX The present sequence is human placental bikunin, which  
XX inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
XX Bikunin can be used to treat or prevent brain and spinal cord  
XX oedema, inflammation, infection or granulomatosis, multiple  
XX sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
XX fibrosis, blood coagulation diseases, polytrauma, stroke,  
XX cerebral or subarachnoid haemorrhage and gastric or cervical

CC cancer and prevent metastasis. It is particularly useful for  
CC reducing blood loss during surgery, and can also be used to treat  
CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
CC influenza and similar viral infections, acute pancreatitis and  
CC gout, and prevent pre-term labour. It has similar properties to  
CC aprotinin, but is less highly charged so should be less  
CC immunogenic and less likely to damage the kidneys. Manipulation  
CC of the bikunin sequence may allow the inhibitory profile to be  
CC altered. It also reduces or eliminates the need for whole donor  
CC blood or blood products during surgery, thereby reducing the risk  
CC of infection and other adverse side effects, as well as reducing  
CC the cost of surgery.

XX Sequence 240 AA;

Query Match 100.0%; Score 170; DB 18; Length 240;  
Best Local Similarity 100.0%; Pred. No. 3.2e-156;  
Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ADERSIHDFCLVSKVVGCRASMPRWYNTDSCOLFVYGGDGNNSNYLTKEECLKK 60  
Db 28 adersihdfclvskvvgcrasmprrwvntdgsclfvvgcdgnsnyltkeecclkk 87

QY 61 CATVTENATGDLATSRNAADSSVPSAPRRQDSEHSDMFNEEYCTANAVTGPCRASFP 120

Db 88 catvtenatgdlatsrnaadssvpsaprrqdsedhssdmfneeyctanavtgcrafp 147

QY 121 RWYFDVERNSCNFIYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK 170

Db 148 rwyfdvernsnfnfiyggcrngknsyrseacmlrcfrqenpplplgsk 197

RESULT 8

W30044

ID W30044 standard; Protein: 248 AA.

XX W30044;

DT 20-APR-1998 (first entry)

DE Human consensus bikunin.

KW Human; consensus bikunin; inhibition; trypsin; kallikrein;  
KW plasmin; factor XIIa; treatment; prevention; oedema;  
KW inflammation; infection; granulomatosis; multiple sclerosis;  
KW ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
KW blood coagulation disease; polytrauma; stroke; haemorrhage;  
KW gastric cancer; cervical cancer; metastasis; blood loss.

OS Homo sapiens.

PN W09733996-A2.

PD 18-SEP-1997.

PF 10-MAR-1997; 97WO-US03894.

PR 04-OCT-1996; 96US-0725251.

PR 11-MAR-1996; 96US-0013106.

PR 14-JUN-1996; 96US-0019793.

PA (FARB ) BAYER CORP.

XX Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;

XX WPI; 1997-470876/43.

XX N-PSDB; T90733.

XX New human placental bikunin - used to inhibit kallikrein, trypsin

XX etc. in treatment of oedema, multiple sclerosis, fibrosis, or

XX perioperative blood loss



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XX 19-FEB-1997.
PD
XX 23-JUL-1996; 96EP-0111861.
PF
XX 24-JUL-1995; 95JP-0187134.
PR
XX (MITU ) MITSUBISHI CHEM CORP.
PA
XX Kawaguchi T, Kitamura N, Shimomura T;
PI
XX WPI; 1997-134770/13.
DR
XX N-PSDB; T61439.
DR
XX Novel protein HAI-II - inhibits protease activity of hepatocyte
PT growth factor activator
XX
XX Claim 4; Page 18-19; 24pp; English.
XX
XX This sequence comprises a novel protein, designated HAI-II,
XX that inhibits the protease activity of hepatocyte growth factor
XX (HGF) activator. The sequence was deduced from a cDNA clone
XX ("T61439") obtained from cancer cell line MKN45. Also claimed
XX are isolated peptides (W13662-64) of HAI-II, the DNA encoding
XX HAI-II, a vector carrying this DNA, and a host cell, pref. an
XX animal cell, transformed with the vector. HAI-II can be used for
XX regulating HGF activator activity (and thus HGF activity) in vitro
XX and in vivo. It may also be used for investigating the function of
XX HAI-II in vivo and the effect of HAI-II in hepatic disorders.
XX
XX Sequence 252 AA;
XX
XX Query Match 100.0%; Score 170; DB 18; Length 252;
XX Best Local Similarity 100.0%; Pred. No. 3.3e-156;
XX Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 ADERSIHDFCLVSKVVGRCRSMRWYNTDGSOLFVYGGDGNNSNYLTKECLKK 60
XX |
XX 28 adersi hdfclvskvvgrcrasmprwvnyntdgsqclfvvgcdgnsnyltkeclkk 87
XX
XX 61 CATVTENATGDLATSRNAADSSVPSAPRRQDSHDHSDMFNYEYCTANAVTGPCRSFP 120
XX |
XX 88 catvt enatgdlatsrnaadssvpsaprrqdsdhdssdmfnyeyctanavtgp rcsfp 147
XX
XX 121 RWFYDVERNSCNFFYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK 170
XX |
XX 148 rwyfdverns cnff yggcr gknksyrseacmlrcfrqenpplplgsk 197
XX
XX RESULT 11
XX W70286
XX ID W70286 standard; Protein; 252 AA.
XX AC
XX W70286;
XX
XX 06-NOV-1998 (first entry)
XX
XX Human tissue factor pathway inhibitor-3 (TFPI-3).
XX
XX Human tissue factor pathway inhibitor-3; TFPI-3; blood clot; sepsis;
XX fibrin clot; coronary occlusion; acute myocardial infarction;
XX prophylaxis; peripheral arterial embolism; inflammatory disease;
XX transplant rejection; anticoagulant; blood transfusion;
XX extracorporeal circulation; dialysis; haemophilia; kunitz type domain.
XX
XX Homo sapiens.
XX
XX OS
XX FH Location/Qualifiers
XX FT Key 1..27
XX FT Peptide /note= "Signal peptide"
XX FT Protein 28..252
XX FT /note= "TFPI-3"
XX FT
XX
XX WO9833920-A2.
XX
XX 06-AUG-1998.
XX
XX 27-JAN-1998; 98WO-US01468.
XX
XX 31-JAN-1997; 97US-0036703.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Gentz RL, Hsu T, Ni J, Rosen CA;
XX
XX WPI; 1998-437473/37.
XX
XX N-PSDB; V33063.
XX
XX Isolated tissue factor pathway inhibitor-3 - used to develop
XX products for treating, e.g. pulmonary embolism, thrombosis, sepsis,
XX inflammatory disease, transplant rejection or haemophilia
XX
XX Disclosure; Fig 1A-1B; 57pp; English.
XX
XX The present sequence represents a human tissue factor pathway
XX inhibitor-3 (TFPI-3) which contains two kunitz type domains. The
XX invention also provides the TFPI-3 cDNA and screening methods for
XX identifying agonists and antagonists of TFPI-3. As TFPI-3 inhibits
XX protease activity, it is claimed to be useful for, e.g. inhibiting
XX intravascular clotting and preventing the formation of fibrin clots
XX both in vitro and in vivo, for treating coronary occlusion with acute
XX myocardial infarction and in the prophylaxis and treatment of
XX peripheral arterial embolism, for the treatment of sepsis, inflammatory
XX diseases and transplant rejection. TFPI-3 is also claimed to be useful
XX as an anticoagulant in blood transfusions, extracorporeal circulation,
XX and dialysis procedures and in blood samples for laboratory purposes.
XX The TFPI-3 antagonists are claimed to be useful for promoting
XX coagulation, e.g. in the treatment of haemophilia.
XX
XX Sequence 252 AA;
XX
XX Query Match 100.0%; Score 170; DB 19; Length 252;
XX Best Local Similarity 100.0%; Pred. No. 3.3e-156;
XX Matches 170; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 ADERSIHDFCLVSKVVGRCRSMRWYNTDGSOLFVYGGDGNNSNYLTKECLKK 60
XX |
XX 28 adersi hdfclvskvvgrcrasmprwvnyntdgsqclfvvgcdgnsnyltkeclkk 87
XX
XX 61 CATVTENATGDLATSRNAADSSVPSAPRRQDSHDHSDMFNYEYCTANAVTGPCRSFP 120
XX |
XX 88 catvt enatgdlatsrnaadssvpsaprrqdsdhdssdmfnyeyctanavtgp rcsfp 147
XX
XX 121 RWFYDVERNSCNFFYGGCRGNKNSYRSEACMLRCFRQENPPLPLGSK 170
XX |
XX 148 rwyfdverns cnff yggcr gknksyrseacmlrcfrqenpplplgsk 197
XX
XX RESULT 12
XX W30051
XX ID W30051 standard; Protein; 153 AA.
XX AC
XX W30051;
XX
XX 20-APR-1998 (first entry)
XX
XX Human placental bikunin.
XX
XX Human; placental bikunin; inhibition; trypsin; kallikrein;
XX plasmin; factor XIIa; treatment; prevention; oedema;
XX inflammation; infection; granulomatosis; multiple sclerosis;
XX ischaemia; perioperative blood loss; sepsis; shock; fibrosis;
XX blood coagulation disease; polytrauma; stroke; haemorrhage;
XX gastric cancer; cervical cancer; metastasis; blood loss.
XX

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XX OS Homo sapiens.
XX PN WO9733996-A2.
XX PD 18-SEP-1997.
XX PF 10-MAR-1997; 97WO-US03894.
XX PR 04-OCT-1996; 96US-0725251.
XX PR 11-MAR-1996; 96US-0013106.
XX PR 14-JUN-1996; 96US-0019793.
XX PA (FARB ) BAYER CORP.
XX PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;
XX DR WPI; 1997-470876/43.
XX PT New human placental bikunin - used to inhibit kallikrein, trypsin
XX PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or
XX PT perioperative blood loss
XX PS Claim 1; Page 67; 110pp; English.
XX CC The present sequence is a human placental bikunin, which
XX CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.
XX CC Bikunin can be used to treat or prevent brain and spinal cord
XX CC oedema, inflammation, infection or granulomatosis, multiple
XX CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,
XX CC fibrosis, blood coagulation diseases, polytrauma, stroke,
XX CC cerebral or subarachnoid haemorrhage and gastric or cervical
XX CC cancer and prevent metastasis. It is particularly useful for
XX CC reducing blood loss during surgery, and can also be used to treat
XX CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,
XX CC influenza and similar viral infections, acute pancreatitis and
XX CC gout, and prevent pre-term labour. It has similar properties to
XX CC aprotinin, but is less highly charged so should be less
XX CC immunogenic and less likely to damage the kidneys. Manipulation
XX CC of the bikunin sequence may allow the inhibitory profile to be
XX CC altered. It also reduces or eliminates the need for whole donor
XX CC blood or blood products during surgery, thereby reducing the risk
XX CC of infection and other adverse side effects, as well as reducing
XX CC the cost of surgery.
XX CC Sequence 153 AA;

Query Match 90.0%; Score 153; DB 18; Length 153;
Best Local Similarity 100.0%; Pred. No. 5.2e-140;
Matches 153; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 IHDFCLVSKVVGCRASMPRWYVNTDGSQCLFYVGGCDGNSNNYLTKEECLKKCATVTE 66
DB 1 ihdfclvskvvgcrasmprrwvntdgsqclfyvggcdgnsnnyltkeec.lkccatvte 60

QY 67 NATGDLATSRNADSSVPSAPRRDSEHSDMFNYEYCTANAVTGPCRASFPFRWYFDV 126
DB 61 natgdlatrsnaadssvpsaprrqdsehdssdmfnyeyctanavtgpccrasfprwfyfdv 120

QY 127 ERNSCNFIYGGCRGNKNSYRSEACMLRCFRQ 159
DB 121 ernscnnfiyggcrgnknsyrseacmlrcfrq 153

RESULT 13
W30052
ID W30052 standard; Protein; 146 AA.
XX AC W30052;
XX DT 20-APR-1998 (first entry)
XX

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DE XX Human placental bikunin.
KW XX Human; placental bikunin; inhibition; trypsin; kallikrein;
KW XX plasmin; factor XIIa; treatment; prevention; oedema;
KW XX inflammation; infection; granulomatosis; multiple sclerosis;
KW XX ischaemia; perioperative blood loss; sepsis; shock; fibrosis;
KW XX blood coagulation disease; polytrauma; stroke; haemorrhage;
KW XX gastric cancer; cervical cancer; metastasis; blood loss.
OS Homo sapiens.
XX PN WO9733996-A2.
XX PD 18-SEP-1997.
XX PF 10-MAR-1997; 97WO-US03894.
XX PR 04-OCT-1996; 96US-0725251.
XX PR 11-MAR-1996; 96US-0013106.
XX PR 14-JUN-1996; 96US-0019793.
XX PA (FARB ) BAYER CORP.
XX PI Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;
XX DR WPI; 1997-470876/43.
XX PT New human placental bikunin - used to inhibit kallikrein, trypsin
XX PT etc. in treatment of oedema, multiple sclerosis, fibrosis, or
XX PT perioperative blood loss
XX PS Claim 1; Page 67; 110pp; English.
XX CC The present sequence is a human placental bikunin, which
XX CC inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.
XX CC Bikunin can be used to treat or prevent brain and spinal cord
XX CC oedema, inflammation, infection or granulomatosis, multiple
XX CC sclerosis, ischaemia, perioperative blood loss, sepsis, shock,
XX CC fibrosis, blood coagulation diseases, polytrauma, stroke,
XX CC cerebral or subarachnoid haemorrhage and gastric or cervical
XX CC cancer and prevent metastasis. It is particularly useful for
XX CC reducing blood loss during surgery, and can also be used to treat
XX CC other cancer, arthritis, anaemia, non-insulin dependent diabetes,
XX CC influenza and similar viral infections, acute pancreatitis and
XX CC gout, and prevent pre-term labour. It has similar properties to
XX CC aprotinin, but is less highly charged so should be less
XX CC immunogenic and less likely to damage the kidneys. Manipulation
XX CC of the bikunin sequence may allow the inhibitory profile to be
XX CC altered. It also reduces or eliminates the need for whole donor
XX CC blood or blood products during surgery, thereby reducing the risk
XX CC of infection and other adverse side effects, as well as reducing
XX CC the cost of surgery.
XX CC Sequence 146 AA;

Query Match 85.9%; Score 146; DB 18; Length 146;
Best Local Similarity 100.0%; Pred. No. 2.7e-133;
Matches 146; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 CLVSKVVGCRASMPRWYVNTDGSQCLFYVGGCDGNSNNYLTKEECLKKCATVTEATG 70
DB 1 clvskvvgcrasmprrwvntdgsqclfyvggcdgnsnnyltkeec.lkccatvtenatg 60

QY 71 DLATSRNADSSVPSAPRRDSEHSDMFNYEYCTANAVTGPCRASFPFRWYFDV 130
DB 61 dlatrsnaadssvpsaprrqdsehdssdmfnyeyctanavtgpccrasfprwfyfdv 120

QY 131 CNNFYGGCRGNKNSYRSEACMLRC 156
DB 121 cnnfiyggcrgnknsyrseacmlrc 146

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||||| catvtenatgdlatsrnaadsvpsaprrqds 92

RESULT 14  
#30054  
ID W30054 standard; Protein; 92 AA.  
XX AC W30054;  
XX DT 20-APR-1998 (first entry)  
XX DE Human placental bikunin.  
XX Human; placental bikunin; inhibition; trypsin; kallikrein;  
XX plasmin; factor XIIa; treatment; prevention; oedema;  
XX inflammation; infection; granulomatosis; multiple sclerosis;  
XX ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
XX blood coagulation disease; polytrauma; stroke; haemorrhage;  
XX gastric cancer; cervical cancer; metastasis; blood loss.  
XX Homo sapiens.  
XX WO9733996-A2.  
XX 18-SEP-1997.  
XX 10-MAR-1997; 97WO-US03894.  
XX 04-OCT-1996; 96US-0725251.  
XX 11-MAR-1996; 96US-0013106.  
XX 14-JUN-1996; 96US-0019793.  
XX (FARB ) BAYER CORP.  
XX Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
XX WPI; 1997-470876/43.  
XX New human placental bikunin - used to inhibit kallikrein, trypsin  
XX etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
XX perioperative blood loss  
XX Claim 1: Page 67; 110pp; English.  
XX The present sequence is a human placental bikunin, which  
XX inhibits, e.g. trypsin, kallikrein, plasmin and factor XIIa.  
XX Bikunin can be used to treat or prevent brain and spinal cord  
XX oedema, inflammation, infection or granulomatosis, multiple  
XX sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
XX fibrosis, blood coagulation diseases, polytrauma, stroke,  
XX cerebral or subarachnoid haemorrhage and gastric or cervical  
XX cancer and prevent metastasis. It is particularly useful for  
XX reducing blood loss during surgery, and can also be used to treat  
XX other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
XX influenza and similar viral infections, acute pancreatitis and  
XX gout, and prevent pre-term labour. It has similar properties to  
XX aprotinin, but is less highly charged so should be less  
XX immunogenic and less likely to damage the kidneys. Manipulation  
XX of the bikunin sequence may allow the inhibitory profile to be  
XX altered. It also reduces or eliminates the need for whole donor  
XX blood or blood products during surgery, thereby reducing the risk  
XX of infection and other adverse side effects, as well as reducing  
XX the cost of surgery.  
XX Sequence 92 AA;  
XX  
XX Query Match 54.1%; Score 92; DB 18; Length 92;  
XX Best Local Similarity 100.0%; Pred. No. 1.9e-81;  
XX Matches 92; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
XX  
XX 1 ADERSIHDFCLVSKVVGCRASPRWYNVTDGSCOLFVYGGDGNNSNYITKECLKK 60  
XX |||||||  
XX 1 adrsihdfclvskvvgcrasprwvnyntdgsqclfyvggdnsgnnyitkeclkk 60  
XX  
XX 61 CATVTENATGDLATSRNADSVPSAPRRQDS 92

Db 61 catvtenatgdlatsrnaadsvpsaprrqds 92  
RESULT 15  
W30063  
ID W30063 standard; Protein; 169 AA.  
XX AC W30063;  
XX DT 20-APR-1998 (first entry)  
XX DE EST R74593 protein.  
XX Human; consensus bikunin; inhibition; trypsin; kallikrein;  
XX plasmin; factor XIIa; treatment; prevention; oedema;  
XX inflammation; infection; granulomatosis; multiple sclerosis;  
XX ischaemia; perioperative blood loss; sepsis; shock; fibrosis;  
XX blood coagulation disease; polytrauma; stroke; haemorrhage;  
XX gastric cancer; cervical cancer; metastasis; blood loss;  
XX EST R74593.  
XX Homo sapiens.  
XX OS  
XX Key Location/Qualifiers  
XX Misc-difference 2 /note= "encoded by TAA"  
XX FT  
XX Misc-difference 23 /note= "encoded by TGA"  
XX FT  
XX Misc-difference 132 /note= "encoded by TGA"  
XX FT  
XX Misc-difference 135 /note= "encoded by TGA"  
XX FT  
XX Misc-difference 160 /note= "encoded by GAN"  
XX FT  
XX Misc-difference 167 /note= "encoded by TGA"  
XX FT  
XX WO9733996-A2.  
XX 18-SEP-1997.  
XX 10-MAR-1997; 97WO-US03894.  
XX 04-OCT-1996; 96US-0725251.  
XX 11-MAR-1996; 96US-0013106.  
XX 14-JUN-1996; 96US-0019793.  
XX (FARB ) BAYER CORP.  
XX Davis G, Delaria KA, Marlor CW, Muller DK, Tamburini PP;  
XX WPI; 1997-470876/43.  
XX N-PSDB; T90736.  
XX New human placental bikunin - used to inhibit kallikrein, trypsin  
XX etc. in treatment of oedema, multiple sclerosis, fibrosis, or  
XX perioperative blood loss  
XX Disclosure; Fig. 2; 110pp; English.  
XX The present sequence is the EST R74593 protein, which is similar  
XX to human bikunin. Bikunin inhibits, e.g. trypsin, kallikrein,  
XX plasmin and factor XIIa.  
XX Bikunin can be used to treat or prevent brain and spinal cord  
XX oedema, inflammation, infection or granulomatosis, multiple  
XX sclerosis, ischaemia, perioperative blood loss, sepsis, shock,  
XX fibrosis, blood coagulation diseases, polytrauma, stroke,  
XX cerebral or subarachnoid haemorrhage and gastric or cervical  
XX cancer and prevent metastasis. It is particularly useful for  
XX reducing blood loss during surgery, and can also be used to treat  
XX other cancer, arthritis, anaemia, non-insulin dependent diabetes,  
XX influenza and similar viral infections, acute pancreatitis and

